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88392

From: Schnizer, Richard  
Sent: Monday, March 10, 2003 12:11 PM  
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Subject: 09/836,439

Please search the commercial databases for SEQ ID NOS: 1-6 from 09/836,439.

Thank you-

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0315

Point of Contact:  
Toby Port  
Technical Info. Specialist  
CM1 6A04  
703-308-3534

Searcher: \_\_\_\_\_  
Phone: \_\_\_\_\_  
Location: \_\_\_\_\_  
Date Picked Up: 3/11  
Date Completed: 3/19  
Searcher Prep/Review: \_\_\_\_\_  
Clerical: \_\_\_\_\_  
Online time: \_\_\_\_\_

TYPE OF SEARCH:

NA Sequences: \_\_\_\_\_  
AA Sequences: \_\_\_\_\_  
Structures: \_\_\_\_\_  
Bibliographic: \_\_\_\_\_  
Litigation: \_\_\_\_\_  
Full text: \_\_\_\_\_  
Patent Family: \_\_\_\_\_  
Other: \_\_\_\_\_

VENDOR/COST (where applic.)

STN: \_\_\_\_\_  
DIALOG: \_\_\_\_\_  
Questel/Orbit: \_\_\_\_\_  
DRLink: \_\_\_\_\_  
Lexis/Nexis: \_\_\_\_\_  
Sequence Sys.: \_\_\_\_\_  
WWW/Internet: \_\_\_\_\_  
Other (specify): \_\_\_\_\_



score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

OM nucleic - nucleic search, using sw model

(without alignments)  
3161.870 Million cell updates/sec

Sequence: 1 ccttccaacctagtagcag.....ggaagggcgcgcttctcgcgc 68

Gapop 10.0 , Gapext 1.0

Total number of hits satisfying chosen parameters: 4109280

Post-processing: Minimum Match 99

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Post-processing:  Minimum Match 0%
                  Maximum Match 100%
                  Listing first 45 summaries
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Database :

1:	gb_ba.*
2:	gb_hlg.*
3:	gb_in.*
4:	gb_om.*
5:	gb_ov.*
6:	gb_pat.*
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12:	gb_sy.*
13:	gb_un.*
14:	gb_vl.*
15:	em_ba.*
16:	em_fun.*
17:	em_hum.*
18:	em_in.*
19:	em_mu.*
20:	em_om.*
21:	em_or.*
22:	em_ov.*
23:	em_pt.*
24:	em_ph.*
25:	em_pl.*
26:	em_ro.*
27:	em_sts.*
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30:	em_hlg_hum.*
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41:	em_hlgo_other.*

Pred. No. is the number of results predicted by chance to have a

Result No.	Score	Query Match	Length	DB	ID	Description
1	28	41.2	68	6	AR024440	AR024440 Sequence
2	28	41.2	68	6	193657	193657 Sequence 23
3	27.2	40.0	10591	3	AC020132	AC020132 Drosophila
4	27.2	40.0	73076	3	AC030355	AC030355 Drosophila
5	27.2	40.0	168540	3	AC092189	AC092189 Drosophila
6	27.2	40.0	303356	3	AE003583	AE003583 Drosophila
7	27	39.7	3554	3	DMFT21	X00854 Drosophila
8	27	39.7	56043	2	AC012761	AC012761 Drosophila
9	27	39.7	66991	3	AC001653	AC001653 Drosophila
10	27	39.7	74026	2	AC101196	AC101196 Mus muscu
11	27	39.7	80866	2	AC012649	AC012649 Drosophila
12	27	39.7	110000	3	AE001572	Confusion (2 of
13	27	39.7	190652	3	AC095015	AC095015 Drosophila
14	27	39.7	208375	2	AC107703	AC107703 Mus muscu
15	27	39.7	309357	3	AE003673	AE003673 Drosophila
16	26.6	39.1	91104	2	AC112364	AC112364 Rattus no
17	26.6	39.1	160589	10	AL663066	AL663066 Mouse DNA
18	26.4	38.8	68	6	AR024439	AR024439 Sequence
19	26.4	38.8	68	6	193655	193655 Sequence 22
20	26.4	38.8	20281	1	AE008861	AE008861 Salmonella
21	26.4	38.8	192675	10	AL589767	AL589767 Mouse DNA
22	26.4	38.8	199842	2	AC094068	AC094068 Rattus no
23	26.4	38.8	207922	10	AL450339	AL450339 Mouse DNA
24	26.2	38.5	139877	9	AC004066	AC004066 Homo sap1
25	26.2	38.5	194521	2	AC093696	AC093696 Homo sap1
26	26	38.2	161119	2	AC109157	AC109157 Mus muscu
27	26	38.2	183311	2	AC109788	AC109788 Bos tauru
28	26	38.2	187352	10	AL607109	AL607109 Mouse DNA
29	26	38.2	222941	2	AC102236	AC102236 Mus muscu
30	25.8	37.9	2693	10	MMPB	X60133 Murine MPB
31	25.8	37.9	2703	10	MMPBALT	X87952 Murine MPB
32	25.8	37.9	2743	10	MMPDE	X55368 Mouse mRNA
33	25.8	37.9	129790	3	AG441131	AJ441131 Anopheles
34	25.8	37.9	157418	2	AC006512	AC006512 Caenorhab
35	25.8	37.9	163475	9	AC092619	AC092619 Homo sap1
36	25.6	37.6	154552	9	AL390964	AL390964 Human DNA
37	25.6	37.6	190721	8	AP003263	AP003263 Oryza sat
38	25.4	37.4	435	6	AX341469	AX341469 Sequence
39	25.4	37.4	1224	9	BC007520	BC007520 Homo sap1
40	25.4	37.4	1324	9	AK026222	AK026222 Homo sap1
41	25.4	37.4	2727	6	AX282833	AX282833 Sequence
42	25.4	37.4	2730	6	AX282845	AX282845 Sequence
43	25.4	37.4	2874	6	AX282837	AX282837 Sequence
44	25.4	37.4	2877	6	AX282849	AX282849 Sequence
45	25.4	37.4	2979	6	AX282841	AX282841 Sequence

RESULT 1				
AR024440				
LOCUS	AR024440	68 bp	DNA	Linear
DEFINITION	Sequence 23 from patent US 5795972.			PAT 05-DEC-1996
ACCESSION	AR024440			
VERSION	AR024440.1	GI:3977734		
KEYWORDS	.			
SOURCE	Unknown.			
ORGANISM	Unknown.			
REFERENCE	Unclassified.			
AUTHORS	1 (bases 1 to 68)			
TITLE	Kmiec, E.B.			
JOURNAL	Chimeric mutational vectors having non-natural nucleotides			
FEATURES	Patent: US 5795972-A 23 18-AUG-1998;			
	Location/Qualifiers			

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source      1. .68
            /organism="unknown"
BASE COUNT      11 a      19 c      19 g      19 t
ORIGIN

Query Match      41.2%; Score 28; DB 6; Length 68;
Best Local Similarity 61.7%; Pred. No. 4.3;
Matches 37; Conservative 3; Mismatches 20; Indels 0; Gaps 0;

OY      9      CCTACGTAGCAGAAAGTTTACUUCUCGACGTAGCGUUGAAGGCGCGTTTCCGCG 68
            ||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
            ||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
            9      CCTGAGGAGAGACTGCTTTGACGCTCTCTCCTCAGAGTCAAGTCCGCGTTTCCGCG 68

RESULT 2
LOCUS      193657      68 bp      DNA      linear      PAT 01-DEC-1998
DEFINITION      Sequence 23 from patent US 5731181.
ACCESSION      193657
VERSION      193657.1 GI:3938127
KEYWORDS
SOURCE      Unknown.
ORGANISM      Unknown.
REFERENCE      1 (bases 1 to 68)
AUTHORS      Kmiec, E.B.
TITLE      Chimeric mutational vectors having non-natural nucleotides
JOURNAL      Patent: US 5731181-A 23-24-MAR-1998;
FEATURES
SOURCE      1. .68
            /organism="unknown"
BASE COUNT      11 a      19 c      19 g      19 t
ORIGIN

Query Match      41.2%; Score 28; DB 6; Length 68;
Best Local Similarity 61.7%; Pred. No. 4.3;
Matches 37; Conservative 3; Mismatches 20; Indels 0; Gaps 0;

OY      9      CCTACGTAGCAGAAAGTTTACUUCUCGACGTAGCGUUGAAGGCGCGTTTCCGCG 68
            ||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
            ||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
            9      CCTGAGGAGAGACTGCTTTGACGCTCTCTCCTCAGAGTCAAGTCCGCGTTTCCGCG 68

RESULT 3
AC020132/c
LOCUS      AC020132      10591 bp      DNA      linear      HTG 03-JAN-2000
DEFINITION      Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in ordered
pieces.
ACCESSION      AC020132
VERSION      AC020132.1 GI:6664765
KEYWORDS      HTG; HTGS; PHASE2.
SOURCE      Drosophila melanogaster.
ORGANISM      Drosophila melanogaster.
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 10591)
Adams, M. and Venter, J.C.
Direct Submission
Submitted (30-DEC-1999) Celera Genomics, 45 West Gude Drive,
Rockville, MD, USA
This sequence was identified as CDN:10212153 by the submitter.
For more information on this record e-mail to fly@celera.com.
* NOTE: This is a 'working draft' sequence.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

FEATURES
source      1. .10591
            /organism="Drosophila melanogaster"
            /db_xref="taxon:7227"
BASE COUNT      2920 a      2216 c      2177 g      3278 t
ORIGIN

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Query Match      40.0%; Score 27.2; DB 2; Length 10591;
Best Local Similarity 60.7%; Pred. No. 12;
Matches 34; Conservative 4; Mismatches 18; Indels 0; Gaps 0;

OY      11      TAGCTAGCAGAAAGTTTACUUCUCGACGTAGCGUUGAAGGCGCGTTTCCGCG 66
            ||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
            ||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
            Db      3352 TAGCTAGCAGAAAGTTTACUUCUCGACGTAGCGUUGAAGGCGCGTTTCCGCG 3297

RESULT 4
LOCUS      AC003055      73076 bp      DNA      linear      INV 04-NOV-1997
DEFINITION      Drosophila melanogaster (PI DS06332 (D91)) DNA sequence, complete
sequence.
ACCESSION      AC003055
VERSION      AC002945 AC002946 AC001988 AC001995 AC002951 AC001991 AC002842
AC001993 AC002948 AC002953 AC001990 AC001989 AC002949 AC002950
AC003055.1 GI:2584828
KEYWORDS      HTG.
SOURCE      Drosophila melanogaster (Subclones in pot2 from PI clone DS06332
(D91)) DNA.
ORGANISM      Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 73076)
Celisner, S.E., Aghavan, A., Arcaina, T.T., Baxter, E., Doyle, C.M.,
Farfan, D.E., Flanagan, J., Houston, K.A., Hummasti, S.R., Karra, K.,
Keane, L., Kim, S.H., Ko, C.L., Li, M., Lomocan, M.A., Mazda, P.,
Mok, M.S., Nixon, R., Pacle, J.M., Park, S., Pfeiffer, B., Punch, D.,
Santos, R.F., Snir, E., Stevko, V., Subramanian, S., Towse, B.,
Wan, K.H., White, K.R., Yee, A., Zhang, R., Zieran, L.L. and
Kimmel, B.
Sequencing of Drosophila chromosome 2L, region 22F
Unpublished (1997)
2 (bases 1 to 73076)
Celisner, S.E., Aghavan, A., Arcaina, T.T., Baxter, E., Doyle, C.M.,
Farfan, D.E., Flanagan, J., Houston, K.A., Hummasti, S.R., Karra, K.,
Keane, L., Kim, S.H., Ko, C.L., Li, M., Lomocan, M.A., Mazda, P.,
Mok, M.S., Nixon, R., Pacle, J.M., Park, S., Pfeiffer, B., Punch, D.,
Santos, R.F., Snir, E., Stevko, V., Subramanian, S., Towse, B.,
Wan, K.H., White, K.R., Yee, A., Zhang, R., Zieran, L.L. and
Kimmel, B.
Direct Submission
Submitted (04-NOV-1997) Berkeley Drosophila Genome Project, MS
74-157, Lawrence Berkeley National Laboratory, One Cyclotron Road,
Berkeley, CA 94720, US
Sequence submitted by:
Lawrence Berkeley National Laboratory, MS 74-157
Berkeley, CA 94720
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive web site (http://www.hgc.lbl.gov/sequence-archive.html) or
send email to drosophila@genome.lbl.gov.
Library location: 66-92.
This PI was assembled from the following subclones: 2_c1
(AC001994), 2_b12 (AC002947), 1_g6 (AC002944), 1_d4 (AC002943),
2_b5, 2_e4 (AC002952), 1_b4, 2_a1, 2_a2, 1_b6 (AC001992), 1_h8
(AC002945), 2_a8 (AC002946), 1_b2 (AC001988), 2_c5 (AC001995),
2_d5 (AC002951), 1_h3 (AC001991), 1_d3 (AC002942), 2_a9 (AC001993),
2_c10 (AC002948), 2_g9 (AC002953), 1_g7 (AC001990), 1_b5
(AC001989), 2_c3 (AC002949), 2_h2, 2_c4 (AC002950), 2_b6.
location/Qualifiers
1. 73076
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/chromosome="2L"
/map="22F2-22F4"
/clone="PI DS06332 (D91)"
BASE COUNT      21203 a      14890 c      15769 g      21214 t
ORIGIN

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/codon_start=1
Query Match      40.0%; Score 27.2; DB 3; Length 303356;
Best Local Similarity 60.7%; Pred. No. 15;
Matches 34; Conservative 4; Mismatches 18; Indels 0; Gaps 0;

OY 11 TAGGTACGAGAAAGTTTTCUUCUGACGTAGUGAGGCGGCTTTTCGC 66
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 277207 TAGCTCGCTGAAGTTTTCCTGCTGCATGTAAATACAGACGANTCTTTCAC 277152

RESULT 7
DMEF21          3554 bp      DNA      linear      INV 07-NOV-1997
LOCUS           Drosophila melanogaster ftz gene.
DEFINITION      X00854 K01951
ACCESSION       X00854.1 GI:7984
VERSION         DNA binding protein; inverted repeat; segmentation gene;
KEYWORDS        unidentified reading frame.
SOURCE          fruit fly.
ORGANISM        Drosophila melanogaster
                Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
                Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
                Ephydroidea; Drosophilidae; Drosophila.
REFERENCE       1 (bases 1 to 3554)
AUTHORS        Laughon, A. and Scott, M. P.
TITLE          Sequence of a Drosophila segmentation gene: protein structure
                homology with DNA-binding proteins
JOURNAL         Nature 310 (5972), 25-31 (1984)
MEDLINE         84245843
PUBMED         6330566
REFERENCE       2 (bases 1 to 3554)
AUTHORS        Scott, M. P. and Weiner, A. J.
TITLE          Structural relationships among genes that control development:
                sequence homology between the Antennapedia, Ultralithorax, and
                fushi tarazu loci of Drosophila
JOURNAL         Proc. Natl. Acad. Sci. U.S.A. 81 (13), 4115-4119 (1984)
MEDLINE         84248068
PUBMED         6330741
COMMENT        Sequence homology to Antp and Ubx genes; see DMANTP1 and DMANTP2
                refer to Pabo C.O., Sauer R.T. (1984) A. Rev. Biochem. 53, 293-321.
                Data kindly reviewed (19-FEB-1986) by A. Laughon.

FEATURES
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                  /db_xref="taxon:7227"
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                  TATA_signal
     TATA_signal 880..896
                  TATA_signal
     pm1_transcript 901..2867
                  pm1_transcript
     exon        901..1777
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     repeat_unit 901..972
                  /note="inverted repeat"
     CDS         924..992
                  /note="unidentified reading frame for pot. minor protein"
                  /codon_start=1
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                  /db_xref="GI:7985"
                  /translation="WQDLRQQLRLKRLTSVACTSOS"
                  979..1048
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                  /protein_id="CAA25408.1"
                  /db_xref="GI:7986"
                  /db_xref="FLYBASE:FBN0001077"
                  /db_xref="SWISS-PROT:P02835"
                  /translation="MATNQSQSHSYSDADNMNMYNMYPHSLPPTYNDNSGNAYONT
                  SNYSYQGYPOESYSESCYYNNQEDVTTQTPVPQPTPPPKATKKRAEDDAASII
                  AAVEBETLRALLNPKYKLKATPDYFTYVEGVKKAPATVTKTASPAASYSQERY
                  TVTPPSAEDVDYLDVYSPOQSTOKLNKNGDPAITPTPTTSLPPLSGISTPPOSFGEK

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SSASVSEINHRIVTAPNGADFNWMSHEETLASDCKDSKRTROTYYRTLEKEF
HENRYTRRRRIDIANALSESEKIKTFORRRKRSKRDLTSSPEHCAGTTAMP
PLEATSTATGASVPVPMHHHOTTAAVPAYSHSHSGGLNDYPOQOTHOODAY
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1066
/note="pot. alternate translation start site for ftz
protein"
intron        1778..1927
                /number=1
exon         1928..2867
                /number=2
misc_feature  1933..2121
                /note="region with homology to Antp and Ubx genes"
                2023..2121
                /note="pot. DNA-binding domain of the ftz protein"
                2041..2043
                /note="GCC (Ala) is GTC (Val) in temperature sensitive
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                2844..2849
                /note="pot. polyadenylation signal"
                2848..2854
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                /note="pot. polyadenylation signal"
BASE COUNT   979 a 952 c 831 g 792 t
ORIGIN
Query Match      39.7%; Score 27; DB 3; Length 3554;
Best Local Similarity 58.8%; Pred. No. 14;
Matches 30; Conservative 6; Mismatches 15; Indels 0; Gaps 0;

OY 15 TAGCAGAAAGTTTTCUUCUGACGTAGUGAGGCGGCTTTTCG 65
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 171 TTGCATTAAGTTTTCCTGCTTACGATTCATTTGGAAAGTCGTTTGTGC 221

RESULT 8
AC012761          56043 bp      DNA      linear      HTG 03-NOV-1999
LOCUS           Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, In ordered
DEFINITION      AC012761
ACCESSION       AC012761.1 GI:6223082
VERSION         AC012761.1
KEYWORDS        HTG; HTGS_PHASE2.
SOURCE          Drosophila melanogaster.
ORGANISM        Drosophila melanogaster
                Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
                Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
                Ephydroidea; Drosophilidae; Drosophila.
REFERENCE       1 (bases 1 to 56043)
AUTHORS        Adams, M. and Venter, J. C.
TITLE          Direct Submission
JOURNAL         Submitted (03-NOV-1999) Celera Genomics, 45 West Gude Drive,
                Rockville, MD, USA
COMMENT        This sequence was identified as CDM:10210082 by the submitter.
                For further information on this sequence you may e-mail to
                fly@celera.com.
                * NOTE: This is a 'working draft' sequence.
                * This sequence will be replaced
                * by the finished sequence as soon as it is available and
                * the accession number will be preserved.

FEATURES
     source      1..56043
                  /organism="Drosophila melanogaster"
                  /db_xref="taxon:7227"
     BASE COUNT  16447 a 11385 c 11452 g 16759 t
     ORIGIN
Query Match      39.7%; Score 27; DB 2; Length 56043;
Best Local Similarity 58.8%; Pred. No. 17;

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Matches 30: Conservative 6: Mismatches 15: Indels 0: Gaps 0:

QY 15 TAGCAGAAAGTTTACUUCGUCACGTAGUGAAGCGCGCTTTTCG 65

Db 5971 TTGCATTAAGTTTACTGTACTAGCATTTTGAAGTGCCTTTGTGG 6021

# RESULT 9

AC001653

66991 bp DNA linear INV 16-APR-1999

# LOCUS

Drosophila melanogaster, chromosome 3R, region 84B1-84B2, P1 clone

# DEFINITION

DS07876, complete sequence.

# ACCESSION

AC001653 L49396 L39779 L32657 L32652 L32645 L32656 L32649 L32648

# VERSION

L32637 L32638 L32639 L32640 L32641 L32646 L32650 L32636 L32643 L32644

# KEYWORDS

HTG.

# SOURCE

Drosophila melanogaster.

# ORGANISM

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

# REFERENCE

1 (bases 1 to 66991)

# AUTHORS

Celniker, S.E., Agbayani, A., Arcaina, T.T., Baxter, E., Blazer, R.G.,

# TITLE

Unpublished (1998)

# JOURNAL

2 (bases 1 to 66991)

# REFERENCE

Martin, C.H., Arcaina, T.T., Bondoc, M.M., Chiang, A., Critz, P.A.,

# AUTHORS

David, C.A., Doyle, C.M., Ericsson, C.L., Farfan, D.E., Gunning, K.M.,

# TITLE

Houston, K.A., Jaklevic, M.A., Kadner, K.E., Kim, K., Kim, S.F.,

# JOURNAL

Ko, C.L., Lewis, K.D., Li, M., Lindquist, K.J., Lomocan, M.A.,

# REFERENCE

Lustre, V.M., Machius, M.U., Mayeda, C.A., Miguel, T.M., Miller, C.A.,

# AUTHORS

Mok, M.S., Pacled, J.M., Patel, S.G., Santos, R.F., Sudamanian, S.,

# TITLE

Wan, K.H., Whitelaw, K.R., Yee, A., Yeh, R.T., Yu, C. and Palazzolo, M.J.

# JOURNAL

Submitted (22-APR-1997) Berkeley Drosophila Genome Project, MS

# COMMENT

64-121, Lawrence Berkeley National Laboratory, One Cyclotron Road,

# FEATURES

Berkeley, CA 94720, US

# Source

On or before Apr 16, 1999 this sequence version replaced g1:483988,

# Location/Qualifiers

g1:483986, g1:1945589, g1:1103946.

# Sequence submitted by:

Berkeley Drosophila Genome Project

# For further information about this sequence, including its location

and relationship to other sequences, please visit our sequence

# archive web site (<http://www.fruitfly.org/sequence/>) or send email

to [bdg@fruitfly.berkeley.edu](mailto:bdg@fruitfly.berkeley.edu).

# P1 library location: 83-4.

1. 66991

# Location/Qualifiers

/organism="Drosophila melanogaster"

# Source

/strain="y2; cn bw sp"

# db\_xref="taxon:7227"

/chromosome="3R"

# map="84B1-84B2"

/clone="PI D507876 (D14)"

# /clone\_id="PI library, partial Sau3a in PMS582tet14ad10"

/note="This sequence has not changed since its original

# submission on 08/25/1997. It was resubmitted in order to

include all secondary accession numbers for the subclones

# belonging to this clone."

BASE COUNT 19565 a 13581 c 13571 g 20274 t

Query Match 39.7% Score 27; DB 3; Length 66991;

Best Local Similarity 58.8%; Pred. No. 17;

Matches 30: Conservative 6: Mismatches 15: Indels 0: Gaps 0:

QY 15 TAGCAGAAAGTTTACUUCGUCACGTAGUGAAGCGCGCTTTTCG 65

Db 25751 TTGCATTAAGTTTACTGTACTAGCATTTTGAAGTGCCTTTGTGG 25801

# RESULT 10

AC101196/c

# LOCUS

Mus musculus clone RP23-177G5, LOW-PASS SEQUENCE SAMPLING.

# DEFINITION

AC101196

# VERSION

AC101196.1 GI:17059970

# KEYWORDS

HTG; HTGS\_PHASE0.

# SOURCE

Mus musculus.

# ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

# REFERENCE

1 (bases 1 to 74026)

# AUTHORS

Birren, B., Linton, L., Nussbaum, C. and Lander, E.

# TITLE

Unpublished

# JOURNAL

2 (bases 1 to 74026)

# REFERENCE

Birren, B., Linton, L., Nussbaum, C., Lander, E., Ali, A., Allen, N.,

# AUTHORS

Anderson, S., Barna, N., Bastien, V., Boguslavsky, L., Boukhalter, B.,

# TITLE

Brown, A., Camarato, J., Campolano, A., Chang, J., Chazaro, B.,

# JOURNAL

Chapel, Y., Colangelo, M., Collins, S., Collamore, A., Cook, A.,

# REFERENCE

Cooke, P., Deatellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S.,

# AUTHORS

Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S.,

# TITLE

Ginde, S., Goid, S., Goyette, M., Graham, L., Grand-Pierre, N.,

# JOURNAL

Hagos, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R.,

# REFERENCE

Jones, C., Kamat, A., Karatas, A., Kells, C., Larocque, K.,

# AUTHORS

Lamazzares, R., Landers, T., Lehoczy, J., Levine, R., Liu, G.,

# TITLE

Maclean, C., McDonald, P., Major, J., Marguis, N., Matthews, C.,

# JOURNAL

McCarthy, M., McEwan, P., McKernan, K., McPheters, R., Meltrin, J.,

# COMMENT

Menus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C.,

# Sequence submitted by:

Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D.,

# For further information about this sequence, including its location

and relationship to other sequences, please visit our sequence

archive web site (<http://www.fruitfly.org/sequence/>) or send email

to [bdg@fruitfly.berkeley.edu](mailto:bdg@fruitfly.berkeley.edu).

1. 74026

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www.seq.wi.mit.edu>

Contact: sequence\_submissions@genome.wi.mit.edu

Center project name: 116139

Center clone name: 177\_G\_5

\* NOTE: This record contains 93 individual

\* sequencing reads that have not been assembled into

\* contigs. Runs of N are used to separate the reads

\* and the order in which they appear is completely

\* arbitrary. Low-pass sequence sampling is useful for

\* identifying clones that may be gene-rich and allows

\* overlap relationships among clones to be deduced.

\* However, it should not be assumed that this clone

\* will be sequenced to completion. In the event that

\* the record is updated, the accession number will

\* be preserved.

1  
706 805: gap of 100 bp in length  
806 1517: contig of 712 bp in length  
1518 1617: gap of 100 bp  
2317 2416: gap of 100 bp  
2417 3114: contig of 698 bp in length  
3115 3214: gap of 100 bp  
3215 3917: contig of 703 bp in length  
3918 4017: gap of 100 bp  
4018 4728: contig of 711 bp in length  
4729 4828: gap of 100 bp  
4829 5529: contig of 701 bp in length  
5530 5629: gap of 100 bp  
5630 6332: contig of 703 bp in length  
6333 6432: gap of 100 bp  
6433 7067: contig of 635 bp in length  
7068 7167: gap of 100 bp  
7168 7877: contig of 710 bp in length  
7878 7977: gap of 100 bp  
7978 8682: contig of 705 bp in length  
8683 8782: gap of 100 bp  
8783 9490: contig of 708 bp in length  
9491 9590: gap of 100 bp  
9591 10300: contig of 710 bp in length  
10301 10400: gap of 100 bp  
10401 11092: contig of 692 bp in length  
11093 11192: gap of 100 bp  
11193 11885: contig of 693 bp in length  
11886 11985: gap of 100 bp  
11986 12684: contig of 699 bp in length  
12685 12784: gap of 100 bp  
12785 13490: contig of 706 bp in length  
13491 13590: gap of 100 bp  
13591 14293: contig of 703 bp in length  
14294 14393: gap of 100 bp  
14394 15099: contig of 706 bp in length  
15100 15199: gap of 100 bp  
15200 15909: contig of 710 bp in length  
15910 16009: gap of 100 bp  
16010 16708: contig of 699 bp in length  
16709 16808: gap of 100 bp  
16809 17515: contig of 707 bp in length  
17516 17615: gap of 100 bp  
17616 18313: contig of 698 bp in length  
18314 18413: gap of 100 bp  
18414 19099: contig of 686 bp in length  
19100 19199: gap of 100 bp  
19200 19896: contig of 697 bp in length  
19897 19996: gap of 100 bp  
19997 20695: contig of 699 bp in length  
20696 20795: gap of 100 bp  
20796 21456: contig of 661 bp in length  
21457 21556: gap of 100 bp  
21557 22217: contig of 661 bp in length  
22218 22317: gap of 100 bp  
22318 23025: contig of 708 bp in length  
23026 23125: gap of 100 bp  
23126 23824: contig of 699 bp in length  
23825 23924: gap of 100 bp  
23925 24610: contig of 686 bp in length  
24611 24710: gap of 100 bp  
24711 25399: contig of 689 bp in length  
25400 25499: gap of 100 bp  
25500 26187: contig of 688 bp in length  
26188 26287: gap of 100 bp  
26288 26993: contig of 706 bp in length  
26994 27093: gap of 100 bp  
27094 27807: contig of 714 bp in length  
27808 27907: gap of 100 bp  
27908 28609: contig of 702 bp in length  
28610 28709: gap of 100 bp  
28710 29413: contig of 704 bp in length

29414 29513: gap of 100 bp  
29514 30190: contig of 677 bp in length  
30191 30290: gap of 100 bp  
30291 30996: contig of 706 bp in length  
30997 31096: gap of 100 bp  
31097 31797: contig of 701 bp in length  
31798 31897: gap of 100 bp  
31898 32567: contig of 670 bp in length  
32568 32667: gap of 100 bp  
32668 33368: contig of 701 bp in length  
33369 33468: gap of 100 bp  
33469 34129: contig of 661 bp in length  
34130 34229: gap of 100 bp  
34230 34935: contig of 706 bp in length  
34936 35035: gap of 100 bp  
35036 35743: contig of 708 bp in length  
35744 35843: gap of 100 bp  
35844 36513: contig of 670 bp in length  
36514 36613: gap of 100 bp  
36614 37323: contig of 710 bp in length  
37324 37423: gap of 100 bp  
37424 38219: contig of 696 bp in length  
38220 38895: contig of 676 bp in length  
38896 38995: gap of 100 bp  
38996 39687: contig of 692 bp in length  
39688 39787: gap of 100 bp  
39788 40501: contig of 714 bp in length  
40502 40601: gap of 100 bp  
40602 41296: contig of 695 bp in length  
41297 41396: gap of 100 bp  
41397 42103: contig of 707 bp in length  
42104 42203: gap of 100 bp  
42204 42906: contig of 703 bp in length  
42907 43006: gap of 100 bp  
43007 43693: contig of 687 bp in length  
43694 43793: gap of 100 bp  
43794 44494: contig of 701 bp in length  
44495 44594: gap of 100 bp  
44595 45291: contig of 697 bp in length  
45292 45391: gap of 100 bp  
45392 46100: contig of 709 bp in length  
46101 46200: gap of 100 bp  
46201 46896: contig of 696 bp in length  
46897 46996: gap of 100 bp  
46997 47682: contig of 686 bp in length  
47683 47782: gap of 100 bp  
47783 48480: contig of 698 bp in length  
48481 48580: gap of 100 bp  
48581 49288: contig of 708 bp in length  
49289 49388: gap of 100 bp  
49389 50070: contig of 682 bp in length  
50071 50170: gap of 100 bp  
50171 50874: contig of 704 bp in length  
50875 50974: gap of 100 bp  
50975 51683: contig of 709 bp in length  
51684 51783: gap of 100 bp  
51784 52495: contig of 712 bp in length  
52496 52595: gap of 100 bp  
52596 53300: contig of 705 bp in length  
53301 53400: gap of 100 bp  
53401 54108: contig of 708 bp in length  
54109 54208: gap of 100 bp  
54209 54904: contig of 696 bp in length  
54905 55004: gap of 100 bp

Query Match 39.7%: Score 27; DB 2; Length 74026;  
Best Local Similarity 67.4%; Pred. No. 17;  
Matches 29; Conservative 4; Mismatches 10; Indels 0; Gaps 0;

QY 21 AAAGTTTACUUCUGCAGUAGGCGGTTT 63  
DB 29154 AAAGTTTACTTTCAGCCAGGCGGTAAGCAGTATT 29112

8487	8565:	gap of unknown length
8567	9246:	contig of 680 bp in length
9247	9326:	gap of unknown length
9327	9911:	contig of 845 bp in length
9972	10051:	gap of unknown length
10052	10632:	contig of 581 bp in length
10633	10712:	gap of unknown length
10713	11338:	contig of 616 bp in length
11329	11408:	gap of unknown length
11409	12070:	contig of 662 bp in length
12071	12150:	gap of unknown length
12151	12954:	contig of 804 bp in length
12955	13034:	gap of unknown length
13035	13884:	contig of 853 bp in length
13888	13967:	gap of unknown length
13968	14675:	contig of 708 bp in length
14676	14755:	gap of unknown length
14756	15777:	contig of 1022 bp in length
15778	15857:	gap of unknown length
15858	16561:	contig of 704 bp in length
16562	16641:	gap of unknown length
16642	17566:	contig of 1015 bp in length
17567	17736:	gap of unknown length
17737	18347:	contig of 611 bp in length
18348	18427:	gap of unknown length
18428	19269:	contig of 842 bp in length
19270	19349:	gap of unknown length
19350	20123:	contig of 774 bp in length
20124	20203:	gap of unknown length
20204	21037:	contig of 834 bp in length
21038	21117:	gap of unknown length
21118	21880:	contig of 763 bp in length
21881	21960:	gap of unknown length
21961	23063:	contig of 1103 bp in length
21962	23143:	gap of unknown length
22064	23742:	contig of 599 bp in length
23144	23882:	gap of unknown length
23745	24697:	contig of 875 bp in length
24698	24777:	gap of unknown length
24778	25699:	contig of 922 bp in length
25700	25779:	gap of unknown length
25780	26547:	contig of 768 bp in length
26548	26637:	gap of unknown length
26638	27682:	contig of 1055 bp in length
27683	27762:	gap of unknown length
27763	28151:	contig of 409 bp in length
28152	28251:	gap of unknown length
28252	28823:	contig of 572 bp in length
28824	28903:	gap of unknown length
28904	29965:	contig of 1062 bp in length
29966	30045:	gap of unknown length
30046	31177:	contig of 1128 bp in length
31174	31253:	gap of unknown length
31254	32156:	contig of 903 bp in length
32157	32236:	gap of unknown length
32237	33872:	contig of 636 bp in length
32873	33952:	gap of unknown length
33953	33817:	contig of 865 bp in length
33818	33897:	gap of unknown length
33898	35066:	contig of 1169 bp in length
35067	35146:	gap of unknown length
35147	37188:	contig of 2042 bp in length
37189	37686:	gap of unknown length
37689	38406:	contig of 1138 bp in length
38407	38486:	gap of unknown length
38487	40075:	contig of 1589 bp in length
40076	41055:	gap of unknown length
41056	41204:	contig of 1049 bp in length
41205	41884:	gap of unknown length
41885	42683:	contig of 1329 bp in length
42684	42694:	gap of unknown length
42694	45330:	contig of 2657 bp in length
45331	45530:	gap of unknown length

```

* 45431 45914: contig of 484 bp in length
* 45915 45994: gap of unknown length
* 45995 45995: contig of 595 bp in length
* 46590 46669: gap of unknown length
* 46670 47222: contig of 553 bp in length
* 47223 47302: gap of unknown length
* 47303 47887: contig of 585 bp in length
* 47888 47968: gap of unknown length
* 47968 48554: contig of 586 bp in length
* 48554 48634: gap of unknown length
* 48634 49267: contig of 633 bp in length
* 49267 49346: gap of unknown length
* 49347 49944: contig of 598 bp in length
* 49945 50024: gap of unknown length
* 50025 50643: contig of 618 bp in length
* 50643 50723: gap of unknown length
* 50723 51238: contig of 505 bp in length
* 51238 51308: gap of unknown length
* 51308 52035: contig of 727 bp in length
* 52035 52115: gap of unknown length
* 52115 52686: contig of 572 bp in length
* 52687 53363: gap of unknown length
* 53363 53443: contig of 597 bp in length
* 53444 53994: gap of unknown length
* 53994 54074: contig of 530 bp in length
* 54074 54435: gap of unknown length
* 54435 55129: contig of 361 bp in length
* 55129 55209: gap of unknown length
* 55210 55887: contig of 615 bp in length
* 55888 55967: gap of unknown length
* 55967 56510: contig of 678 bp in length
* 56510 56590: gap of unknown length
* 56591 57022: contig of 543 bp in length
* 57022 57103: gap of unknown length
* 57103 57394: contig of 432 bp in length
* 57395 57474: gap of unknown length
* 57475 58107: contig of 292 bp in length
* 58107 58187: gap of unknown length
* 58187 58920: contig of 632 bp in length
* 58921 59000: gap of unknown length
* 59001 59609: contig of 734 bp in length
* 59610 59689: gap of unknown length
* 59689 60372: contig of 609 bp in length
* 60372 60451: gap of unknown length
* 60452 61001: contig of 682 bp in length
* 61001 61081: gap of unknown length
* 61082 61656: contig of 550 bp in length
* 61657 61736: gap of unknown length
* 61737 62261: contig of 575 bp in length
* 62262 62341: gap of unknown length
* 62342 62865: contig of 525 bp in length
* 62866 62945: gap of unknown length
* 62946 63538: contig of 524 bp in length
* 63539 63618: gap of unknown length
* 63619 64136: contig of 593 bp in length
* 64137 64216: gap of unknown length

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Query Match
Best Local Similarity 39.7%; Score 27; DB 2; Length 80866;
Matches 30; Conservative 6; Mismatches 15; Indels 0; Gaps 0;
Oy 15 TAGCAGAAAGTTTACUUCUGACAGTACGAGGCGCGTTTTCG 65
Db 25511 TTGCATAAAGTTTACTGTTTACTAGTCAATTTGGAAGTGGCTTTGG 25561

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RESULT 12
AE001572-1/c
WPCOMMENT
Sequence split into 5 fragments LOCUS AE001572 Accession AE001572
Fragment Name Begin End

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AE001572-0 1 110000
AE001572-1 100001 210000
AE001572-2 200001 310000
AE001572-3 300001 410000
AE001572-4 400001 429825
Continuation (2 of 5) of AE001572 from base 100001 (AE001572 Drosophila melanogaster)

```

```

Query Match
Best Local Similarity 39.7%; Score 27; DB 3; Length 110000;
Matches 30; Conservative 6; Mismatches 15; Indels 0; Gaps 0;

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Oy 15 TAGCAGAAAGTTTACUUCUGACAGTACGAGGCGCGTTTTCG 65
Db 93289 TTGCATAAAGTTTACTGTTTACTAGTCAATTTGGAAGTGGCTTTGG 93239

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RESULT 13
AC095015 190642 bp DNA linear INV 15-SEP-2001
DEFINITION Drosophila melanogaster, chromosome 3R, region 84A-84B, BAC clone
LOCUS BACR32J03, complete sequence.
AC095015
VERSION AC095015
KEYWORDS AC095015.1 GI:15624857
SOURCE HTG.
ORGANISM Drosophila melanogaster.
Drosophila melanogaster.
Neoptera: Endopterygota; Diptera: Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 190642)

```

```

REFERENCE
AUTHORS

```

```

TITLE
JOURNAL
REFERENCE
AUTHORS

```

```

2 (bases 1 to 190642)
Celiker, S.E., Adams, M.D., Krommiller, B., Tyler, D., Wan, K.H.,
Holt, R.A., Evans, C.A., Gocayne, J.D., Amanatides, P.G., Brandon, R.C.,
Rogers, Y., An, H., Baldwin, D., Banazon, J., Beeson, K.Y., Busam, D.A.,
Carlson, J.W., Center, A., Champs, M., Davenport, L.B., Dietz, S.M.,
Dodson, K., Dorsett, V., Dou, L.E., Doyle, C., Dresner, D., Farfan, D.,
Fierler, S., Frise, E., Galle, R.F., Garg, N.S., George, R.A.,
Gonzalez, M., Houck, J., Hoskins, R.A., Hostin, D., Howland, T.J.,
Ibegwam, C., Jalali, M., Kruse, D., Li, P., Mettel, B., Moshrefi, A.,
McIntosh, T.C., Moy, M., Murphy, B., Nelson, C., Nelson, K.A., Nunoo, J.,
Pachet, J., Paragas, V., Park, S., Patel, S., Pfeiffer, B.,
Phouanavong, S., Pittman, G.S., Puri, V., Richards, S., Scheeler, F.,
Stapleton, M., Strong, R., Svirskaas, R., Tector, C., Williams, S.M.,
Zaveri, J.S., Smith, H.O., Rubin, G.M. and Venter, J.C.
Sequencing of Drosophila chromosome 3R, region 84A-84B
Unpublished

```

```

TITLE
JOURNAL
COMMENT
Submitted (15-SEP-2001) Berkeley Drosophila Genome Project, MS
64-121, Lawrence Berkeley National Laboratory, One Cyclotron Road,
Berkeley, CA 94720, US
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720

```

This sequence was assembled using end sequences from a whole genome shotgun and from subclones of this BAC and its neighboring clones. For further information about this sequence, including its location and relationship to other sequences, please visit our sequence archive web site (<http://www.fruitfly.org/sequence/>) or send email to [bdg@fruitfly.berkeley.edu](mailto:bdg@fruitfly.berkeley.edu).

FEATURES  
Source Location/Qualifiers  
1.190642  
/organism="Drosophila melanogaster"  
/strain="y; cn bw sp"  
/db\_xref="taxon:7227"  
/chromosome="3R"  
/map="84A-84B"  
/clone="BACR32J03 (D1350)"  
/clone\_lib="RPC1-98 (Roswell Park Cancer Institute  
Drosophila melanogaster BAC library, partial Ecoli in  
pBACE3.6)"

BASE COUNT 55773 a 39290 c 38978 g 56601 t

ORIGIN  
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Best Local Similarity 58.8%; Pred. No. 18;  
Matches 30; Conservative 6; Mismatches 15; Indels 0; Gaps 0;

OY 15 TAGCAGAAAGTTTACUUCUGACGACGAGGAGGCGGCTTTCG 65  
Db 30503 TTGCATTAAGTTTACTGTTACTAGTCATTTGGAAGTGCCTTGTGG 30553

RESULT 14  
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LOCUS  
DEFINITION Mus musculus clone RP23-472H1, WORKING DRAFT SEQUENCE, 8 ordered  
pieces.  
AC107703  
AC107703.5 GI:22381033  
HTG: HTGS\_PHASE2; HTGS\_DRAFT; HTGS\_FUZZTOP.  
KEYWORDS house mouse.  
SOURCE  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 208375)  
Birren, B., Nusbaum, C. and Lander, E.  
Mus musculus, clone RP23-472H1  
Unpublished  
2 (bases 1 to 208375)  
Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,  
Anderson, S., Barina, N., Bastien, V., Boguslavsky, L., Boukhaltier, B.,  
Brown, A., Camarata, J., Campolano, A., Chang, J., Chazaro, B.,  
Choepe, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A.,  
Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Fero, S.,  
Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S.,  
Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,  
Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,  
Kamat, A., Karatas, A., Kells, C., Laroque, K., Lamazares, R.,  
Landers, T., Lehoczy, J., Levine, R., Liu, G., Maclean, C.,  
Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M.,  
McKen, P., McKernan, K., Meldrum, J., Meneus, L., Mihova, T.,  
Mien, P., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C.,  
Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J.,  
Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C.,  
Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J.,  
Rosetti, M., Roy, A., Santos, R., Schauer, S., Schuback, R., Seaman, S.,  
Severy, P., Spencer, B., Strange-Thomann, N., Stojanovic, N.,  
Strauss, N., Sudramanlian, A., Talamas, J., Testafaye, S., Theodore, J.,  
Topham, K., Travers, M., Travis, N., Triggillo, O., Vassiliev, H.,  
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G.,  
Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE  
JOURNAL Submitted (24-JAN-2002) Whitehead Institute/MIT Center for Genome  
Research, 320 Charles Street, Cambridge, MA 02141, USA  
3 (bases 1 to 208375)  
Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,  
Barina, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhaltier, B.,  
Camarata, J., Chang, J., Chazaro, B., Choepe, Y., Collymore, A.,  
Cooke, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S.,  
Fero, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J.,  
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B.,  
Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A.,

FEATURES  
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/db\_xref="taxon:10090"  
/clone="RP23-472H1"  
/clone\_lib="RPC1-23 female Mouse BAC"  
1.53805  
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vector\_side:left"  
misc\_feature  
misc\_feature  
misc\_feature  
53906.56169

NOTE: This is a 'working draft' sequence. It currently  
\* consists of 8 contigs. Gaps between the contigs  
\* are represented as runs of N. The order of the pieces  
\* is believed to be correct as given, however the sizes  
\* of the gaps between them are based on estimates that have  
\* provided by the submitter.  
\* This sequence will be replaced  
\* by the finished sequence as soon as it is available and  
\* the accession number will be preserved.  
1 53805: contig of 53805 bp in length  
53806 53905: gap of 100 bp  
53906 56169: contig of 2264 bp in length  
56170 56269: gap of 100 bp  
56270 62030: contig of 5761 bp in length  
62031 62130: gap of 100 bp  
62131 71647: contig of 9517 bp in length  
71648 71747: gap of 100 bp  
71748 84166: contig of 12419 bp in length  
84167 84266: gap of 100 bp  
84267 138240: contig of 53974 bp in length  
138241 138340: gap of 100 bp  
138341 186299: contig of 47959 bp in length  
186300 186399: gap of 100 bp  
186400 208375: contig of 21976 bp in length.

Center: Whitehead Institute/ MIT Center for Genome Research  
Center code: WIRB  
Web site: <http://www-seq.wi.mit.edu>  
Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)  
Project Information  
Center project name: L18630  
Center clone name: 472\_H-1  
Summary Statistics  
Sequencing vector: plasmid; n/a; 100% of reads  
Chemistry: dye-terminator Big Dye; 100% of reads  
Assembly program: Phrap; version 0.960731  
Consensus quality: 206755 bases at least Q40  
Consensus quality: 207261 bases at least Q30  
Consensus quality: 207461 bases at least Q20  
Insert size: 19400; agarose-fp  
Insert size: 207675; sum-of-contigs  
Quality coverage: 11.1 in Q20 bases; sum-of-contigs  
Quality coverage: 10.4 in Q20 bases; sum-of-contigs



TITLE  
 JOURNAL  
 MEDLINE  
 PUBMED  
 REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL  
 COMMENT  
 FEATURES  
 source  
 gene  
 mRNA  
 CDS  
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 gene  
 mRNA  
 CDS  
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Venter,E., Wang,A.H., Wang,X., Wang,Z.Y., Wasserman,D.A.,  
 Weinster,G.M., Weissenbach,J., Williams,S.M., Woodage,T.,  
 Worley,K.C., Wu,D., Yang,S., Yao,Q.A., Ye,J., Yin,R.F.,  
 Zaveri,I.S., Zhang,M., Zhou,G., Zhao,Q., Zheng,L., Zheng,X.H.,  
 Zhong,F.N., Zhong,W., Zhou,X., Zhu,S., Zhu,X., Smith,H.O.,  
 Gibbs,R.A., Myers,E.W., Rubin,G.M. and Venter,J.C.  
 The genome sequence of Drosophila melanogaster  
 Science 287 (5461), 2185-2195 (2000)  
 20196006  
 10731132  
 2 (bases 1 to 309357)  
 Adams,M.D., Celniker,S.E., Gibbs,R.A., Rubin,G.M. and Venter,C.J.  
 Direct Submission  
 Submitted (21-MAR-2000) Celera Genomics, 45 West Gude Drive,  
 Rockville, MD, USA  
 On Oct 9, 2000 this sequence version replaced gi:7298860.  
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LOANISCTCYADVTPTEGSGGGSSSSANNNNNSANNNSDGLSDIRISK
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gene

**mRNA**

CDS

gene

miRNA

CDS

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```

```

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Matches 30; Conservative 6; Mismatches 15; Indels 0; Gaps 0;

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Search completed: March 17, 2003, 11:24:06
Job time : 920.892 secs

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Search completed: March 17, 2003, 11:24:06  
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XX	PN	M09741141-AI.	
XX	PD	06-NOV-1997.	
XX	PF	01-MAY-1997;	97WO-US07362.
XX	PR	01-MAY-1996;	96US-0640517.
XX	PA	(UYJE-) UNIV JEFFERSON THOMAS.	
XX	PI	Cole-strauss A, Kmiec EB, Yoon K;	
XX	DR	WPI; 1997-549675/50.	
XX	PT	Chimeric nucleic acid repair vectors - used for treating diseases such as sickle cell disease, beta-thalassemia, Gaucher disease,	
XX	PT	hypercholesterolaemia, emphysema or haemophilia	
PS		Disclosure; Fig 3; 79pp: English.	
XX	CC	This chimeric repair vector (CRV) SC4 is used in comparative studies on the experimental use of a CRV SCI designed to repair the mutation found in sickle cell disease beta-globin and the beta-globin of a HSC. The CRV is designed to repair the mutation contains a nucleic acid having at most one 3' end and one 5' end comprising a segment of unpaired bases disposed. The unpaired bases separate the nucleic acid into a first strand and a second strand, comprising a first region and a second region respectively, each region having at least 15 nucleotides. Each nucleotide of the first region is Watson-Crick paired to a nucleotide of the second region and the first region comprises at least 8 ribonucleotides, which are Watson-Crick paired to 2'-deoxynucleotides, which ribonucleotides form at least one ribonucleotide segment of at least 3 ribonucleotides and the sequence of the first or the second region is the sequence of a fragment of a wild-type allele of a human gene. The CRVs can be used for repairing genetic mutations in cells for re-introducing into a patient for treating diseases. They can be used for treating, sickle cell disease, beta-thalassemia, familial hypercholesterolaemia, Gaucher disease, emphysema or haemophilia.	
CC	SQ	Sequence 68 BP; 11 A; 19 C; 19 G; 13 T; 6 U; 0 other;	
CC		Query Match	41.2%; Score 28; DB 18; Length 68;
CC		Best Local Similarity	66.7%; Pred. No. 0.57;
CC		Matches	40; Conservative 0; Mismatches 20; Indels 0; Gaps 0;
OY		9 CCTACGAGCAGAAGATTTCATCUUUCUGACGTAGGUGUGAAGCGCGTTTGCGGC 68   Db 9 CTCGAGGAGAGAACTCTTTCGCAUCUUCUCCCTCAGAGCAGCAGGUGCGGCTTTTCGCCG 68	
RESULT 2			
AAVI2904	DD	AAVI2904 standard; DNA; 68 BP.	
XX	AC	AAVI2904;	
XX	DT	17-JUN-1998 (first entry)	
XX	DE	Chimeric mutational vector SC4.	
KM		Chimeric mutational vector; alkaline phosphatase gene; gene repair;	
KM		thalassaemia-related mutation; human; Gaucher's disease; sickle-cell anaemia;	
KM		thalassaemia; familial hypercholesterolaemia; emphysema; therapy;	
KM		circular; ss.	
OS		Synthetic.	
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FT	/tag= c	
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FT	/tag= f	
XX	/note= "binds to nucleotides 59 to 55"	
PN	M09748714-A1.	
XX	24-DEC-1997.	
PD		
XX	16-JUN-1997;	97MO-US10538.
PF		
XX	17-JUN-1996;	96US-0664487.
PR		
XX	(UYJE-) UNIV JEFFERSON THOMAS.	
PA		
XX	Kmlac EB;	
PI		
XX	WPI; 1998-063068/06.	
DR		
XX	Oligonucleotide for altering a genomic sequence in eukaryotes - particularly for correcting disease-related mutation(s) and for production of transgenic animals and plants	
PT		
XX		
XX	Example 7.2; Fig 3; 68bp; English.	
PS		
XX	This sequence represents a nucleotide analogue of the invention, termed a chimeric mutational vector (CMV). This sequence is directed against the beta-globin gene. The CMVs (I) are for altering a gene in a eukaryotic cell, and comprise: (a) a first strand (S1) having at least 15 nucleotides (nt); at least 3 nuclease-resistant ribo-type nt (nr') and at least 3 contiguous ribo-type nt, the same as, or additional to, the nr'; and (b) a second strand (S2) in which the nt are Watson-Crick (WC) paired to the nt in S1. The contiguous ribo-type nt in S1 are WC-paired to 2'-deoxyribo-type nt, and at least one ribo-type nt is other than a 2'-O-methyl substituted nt. nr' are 2'-AX-nucleosides; 2'-AX-nucleosides or 2'-AR-nucleotides; A = oxygen, fluoro, chloro or bromo; when A = O, X = hydrogen or 1-6C alky1 and R = 1-6C alky1; when A is halo then R and X are absent. (I) are used to repair a disease-related mutation in human cells (e.g. those associated with Gaucher's disease, sickle-cell anaemia, thalassemia, familial hypercholesterolaemia, emphysema etc.). The chimeric mutation vectors are also used to inactivate specific genes, i.e. to generate transgenic ('knockout') animals or plants. They are also CC used for biomedical research and for pharmaceutical production. Any CC eukaryotic gene of known sequence can be altered, by replacement, deletion or addition.	
CC		
XX	Sequence 68 BP; 11 A; 19 C; 19 G; 13 T; 6 U; 0 other:	
SQ		
Query Match	41.2%; Score 28; DB 19; Length 68;	
Best Local Similarity	66.7%; Pred. No. 0.57;	
Matches	40; Conservative 0; Mismatches 20; Indels 0; Gaps 0	
DG		
YY	9 CCTACGTAGCAGAAGATTTTTACUUCUCGCACTAGCGTUGGAAGGGCGGCTTTGCGCC 68   Dbg 9 C CTGAGAGAGAAGACTCTTTTCAGUCUUCUCCCTAGAGACGACGAGUCGCGGTTTTGCGCC 68	
RESULT 3		
AAAX19657		
ID	AAAX19657 standard; DNA; 68 BP.	
XX		
AC	AAAX19657;	
XX		
TJ	02-JUN-1999 (first entry)	

XX	Oligonucleotide SEQ ID NO:96.
DE	
KW	Genome; genetic lesion; haematopoietic stem cell; hepatocyte; RNase;
KW	human wild-type allele; mutation; sickle cell anaemia; thalassemia;
KM	Gaucher's disease; glucocerebrosidase gene; hypercholesterolaemia;
KW	emphysema; haemophilia; Christmas disease; ss.
OS	Synthetic.
XX	
PN	US588993-A.
XX	
PD	30-MAR-1999.
XX	
PF	05-AUG-1997; 97US-0906265.
XX	
PR	05-AUG-1997; 97US-0906265.
PR	01-JAN-1996; 96US-0640517.
XX	17-JUN-1996; 96US-0664487.
PA	(UYJE-) UNTV JEFFERSON THOMAS.
PI	Cole-Strauss AD, Kmiec EB;
DR	WPI; 1999-243264/20.
PT	Double-stranded oligonucleotides with containing a human wild-type
PT	allele - useful for repairing mutations in human cells,
PS	particularly those causing sickle cell anaemia or thalassemia
XX	Disclosure; Column 61-62; 40pp; English.
CC	The present invention describes double-stranded oligonucleotides (I)
CC	containing fragments of wild-type human alleles. (I) are used to repair
CC	disease associated mutations in human cells. (I) are preferably used to
CC	treat sickle cell anaemia or thalassemia (mutations in the beta globin
CC	gene, including the promoter region), or Gaucher's disease (mutations in
CC	the glucocerebrosidase gene), in haematopoietic cells. (I) may also be
CC	used to treat familial hypercholesterolemia (mutations in the low-
CC	density lipoprotein receptor gene), emphysema (the alpha 1-antitrypsin
CC	gene), haemophilias (the factor VIII gene) or Christmas disease (the
CC	factor IX gene), in hepatocytic cells. (I) provides repair of small
CC	genetic mutations. The present sequence represents an oligonucleotide
CC	from the present invention.
XX	
XQ	Sequence 68 BP; 11 A; 19 C; 19 G; 13 T; 6 U; 0 other;
Query Match	41.2%; Score 28; DB 20; Length 68;
Best Local Similarity	66.7%; Pred. NO. 0.57;
Matches 40; Conservative 0; Mismatches 20; Indels 0; Gaps 0;	
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RESULT 4	
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ID	AAVS2984 standard; DNA; 68 BP.
XX	
AAVS2984;	
XX	
DT	04-DEC-1998 (first entry)
DE	Oligonucleotide used in the course of the invention.
XX	
KW	Chimeric oligonucleotide; specific alteration; target sequence;
KW	intramolecular duplex stability; intermolecular duplex stability;
KW	nuclease degradation; chemical stability; hydrolysis resistance;
KW	degradation resistance; A-type helix formation; stable conformation;
SS	DNA/RNA hybrid; cyclic; circular; ss.
SY	Synthetic.

[illegible]

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XX 27-SEP-2001.
PD
XX 23-MAR-2001; 2001MO-US09231.
XX PF
XX 23-MAR-2000; 2000US-191637P.
XX PR
XX 11-JUL-2000; 2000US-0614150.
XX PA
XX (PEKE ) PE CORP NY.
XX PI
XX Venter JC, Adams M, Li PWD, Myers EW;
XX WPI; 2001-656860/75.
XX DR
XX P-PSDB; ABB59865.
XX PT
XX PT New isolated nucleic acid detection reagent for detecting 1000 or more
XX PT genes from Drosophila and for elucidating cell signalling and cell-cell
XX PT interactions -
XX PS
XX Claim 1; SEQ ID NO 6386; 21pp + Sequence Listing; English.
XX CC
XX The invention relates to an isolated nucleic acid detection reagent
XX CC capable of detecting 1000 or more genes from Drosophila. The invention is
XX CC useful in developmental biology and in elucidating cell signalling and
XX CC cell-cell interactions in higher eukaryotes for the development of
XX CC insecticides, therapeutics and pharmaceutical drugs. The invention
XX CC discloses genomic DNA sequences (AB16176-AB130511), expressed DNA
XX CC sequences (AB101840-AB16175) and the encoded proteins
XX CC (ABB5737-ABB72072).
XX CC The sequence data for this patent did not form part of the printed
XX CC specification, but was obtained in electronic format directly from WIPO
XX CC at ftp.wipo.int/pub/published_pct_sequences.
XX SQ
XX Sequence 3785 BP; 876 A; 851 C; 982 G; 1076 T; 0 other;
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XX Query Match 39.7%; Score 27; DB 23; Length 3785;
XX Best Local Similarity 58.8%; Pred. No. 3.3;
XX Matches 30; Conservative 6; Mismatches 15; Indels 0; Gaps 0;
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XX AC
XX AAV09395;
XX XX
XX 14-MAY-1998 (first entry)
XX DT
XX XX
XX Chimeric repair vector (CRV) SC3.
XX DE
XX XX
XX Chimeric repair vector; CRV; treatment; genetic mutation; repair;
XX KW sickle cell disease; beta-thalassemia; Gaucher disease; emphysema;
XX KM hypercholesterolaemia; haemophilia; DNA/RNA hybrid; ss.
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XX Synthetic.
XX OS
XX Homo sapiens.
XX XX
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XX FT /note= "RNA nucleotides"
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XX PN
XX WO9741141-A1.
XX XX
XX 06-NOV-1997.
XX PD
XX XX
XX 01-MAY-1997; 97WO-US07362.
XX PF
XX XX
XX 01-MAY-1996; 96US-0640517.
XX PR
XX XX
XX (UYJE-) UNIV JEFFERSON THOMAS.
XX PA
XX XX
XX Cole-strauss A, Kniec EB, Yoon K;
XX PI
XX WPI; 1997-549675/50.
XX DR
XX XX
XX Chimeric nucleic acid repair vectors - used for treating diseases
XX PT such as sickle cell disease, beta-thalassemia, Gaucher disease,
XX PT hypercholesterolaemia, emphysema or haemophilia
XX PS
XX Disclosure; Fig 3; 79pp; English.
XX XX
XX This chimeric repair vector (CRV) SC3 is used in comparative studies on
XX CC the experimental use of a CRV SC1 designed to repair the mutation found
XX CC in sickle cell disease beta-globin and the beta-globin of a HSC. The CRV
XX CC is designed to repair the mutation contains a nucleic acid having at most
XX CC one 3' end and one 5' end comprising a segment of unpaired bases
XX CC disposed. The unpaired bases separate the nucleic acid into a first
XX CC strand and a second strand, comprising a first region and a second region
XX CC respectively, each region having at least 15 nucleotides. Each nucleotide
XX CC of the first region is Watson-Crick paired to a nucleotide of the second
XX CC region and the first region comprises at least 8 ribonucleotides, which
XX CC are Watson-Crick paired to 2'-deoxynucleotides, which ribonucleotides
XX CC form at least one ribonucleotide segment of at least 3 ribonucleotides
XX CC and the sequence of the first or the second region is the sequence of a
XX CC fragment of a wild-type allele of a human gene. The CRVs can be used for
XX CC repairing genetic mutations in cells for re-introducing into a patient
XX CC for treating diseases. They can be used for treating, sickle cell
XX CC disease, beta-thalassemia, familial hypercholesterolaemia, Gaucher
XX CC disease, emphysema or haemophilia.
XX XX
XX SQ
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XX Query Match 38.8%; Score 26.4; DB 18; Length 68;
XX Best Local Similarity 65.0%; Pred. No. 2.2;
XX Matches 39; Conservative 0; Mismatches 21; Indels 0; Gaps 0;
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XX 9 CCTACGTAGCAGAAAGTTTACUUCUGCAGTACGUGGAGGCGCGTTTCGCGC 68
XX DB 9 CCTAGAGAGAAAGTCTGTTTGCAGCUCUUCCTCAGAGACUAGGCGGTTTTCGCGC 68
XX
XX RESULT 7
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XX ID AAV12903 standard; DNA; 68 BP.
XX AC
XX AAV12903;
XX XX
XX 17-JUN-1998 (first entry)
XX DT
XX XX
XX Chimeric mutational vector SC3.
XX DE
XX XX
XX Chimeric mutational vector; alkaline phosphatase gene; gene repair;
XX KW disease-related mutation; human; Gaucher's disease; sickle-cell anemia;
XX KM thalassemia; familial hypercholesterolaemia; emphysema; therapy;
XX XX
XX Synthetic.
XX OS
XX Key
XX FH Location/Qualifiers
XX FT

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 FT /note= "binds to nucleotides 54 to 30"  
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 FT /note= "binds to nucleotides 25 to 1"  
 FT misc-RNA 30..39  
 FT /tag= c  
 FT /note= "binds to nucleotides 25 to 1"  
 FT misc-RNA 45..54  
 FT /tag= d  
 FT /note= "binds to nucleotides 59 to 55"  
 FT misc-feature 55..59  
 FT /tag= e  
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 FT /tag= f  
 FT /note= "binds to nucleotides 59 to 55"  
 PN WO9748714-A1.  
 XX  
 XX 24-DEC-1997.  
 PD  
 XX  
 XX 16-JUN-1997; 97WO-US10538.  
 PF  
 XX  
 XX 17-JUN-1996; 96US-0664487.  
 PR  
 XX  
 XX (UYJE-) UNIV JEFFERSON THOMAS.  
 PA  
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 XX Kmlec EB:  
 PI  
 XX  
 XX WPI: 1998-063068/06.  
 DR  
 XX  
 XX Oligonucleotide for altering a genomic sequence in eukaryotes -  
 PT particularly for correcting disease-related mutation(s) and for  
 PT production of transgenic animals and plants  
 PS  
 XX Example 7.2; Fig 3; 68bp; English.  
 XX  
 XX This sequence represents a nucleotide analogue of the invention, termed a  
 CC chimeric mutational vector (CMV). This sequence is directed against the  
 CC beta-globin gene. The CMVs (I) are for altering a gene in a  
 CC eukaryotic cell, and comprise: (a) a first strand (S1) having at least 15  
 CC nucleotides (nt); at least 3 nucleotide-resistant ribo-type nt (nt') and at  
 CC least 3 contiguous ribo-type nt, the same as, or additional to, the nt';  
 CC and (b) a second strand (S2) in which the nt are Watson-Crick (WC) paired  
 CC to the nt in S1. The contiguous ribo-type nt in S1 are WC-paired to  
 CC 2'-deoxyribo-type nt, and at least one ribo-type nt is other than a  
 CC 2'-O-methyl substituted nt. nt' are 2'-AX-nucleosides; 2'-AX-nucleosides  
 CC or 2'-AR-nucleosides; A = oxygen, fluoro, chloro or bromo; when A = O,  
 CC X = hydrogen or 1-6C alkyl and R = 1-6C alkyl; when A is halo then R and  
 CC X are absent. (I) are used to repair a disease-related mutation in human  
 CC cells (e.g. those associated with Gaucher's disease, sickle-cell anaemia,  
 CC thalassemia, familial hypercholesterolaemia, emphysema etc.). The  
 CC chimeric mutation vectors are also used to inactivate specific genes,  
 CC i.e. to generate transgenic ('knockout') animals or plants. They are also  
 CC used for biomedical research and for pharmaceutical production. Any  
 CC eukaryotic gene of known sequence can be altered, by replacement,  
 CC deletion or addition.  
 XX  
 XX Sequence 68 BP; 12 A; 18 C; 18 G; 14 T; 6 U; 0 other;  
 SQ  
 Query Match 38.8%; Score 26.4; DB 19; Length 68;  
 Best Local Similarity 65.0%; Pred. No. 2.2;  
 Matches 39; Conservative 0; Mismatches 21; Indels 0; Gaps 0;  
 QY 9 CCTACGTACGAGAAAGTTTACUUCUGCAGTACGAGUUGGAGCGCGTTTGGCGC 68  
 DB 9 CCTGAGAGAGAAAGCTGCTTTGAGUCUUCUCCCTCAGAGAGUAGUAGCGCGTTTGGCGC 68

AC AAX19656;  
 XX  
 XX 02-JUN-1999 (first entry)  
 DT  
 XX  
 XX Oligonucleotide SEQ ID NO:95.  
 DE  
 XX  
 XX Genome; genetic lesion; haematopoietic stem cell; hepatocyte; RNase;  
 KW human wild-type allele; mutation; sickle cell anaemia; thalassemia;  
 KW Gaucher's disease; glucocerebrosidase gene; hypercholesterolaemia;  
 KW emphysema; haemophilia; Christmas disease; ss.  
 XX  
 XX Synthetic.  
 OS  
 XX  
 XX US5888983-A.  
 PN  
 XX  
 XX 30-MAR-1999.  
 PD  
 XX  
 XX 05-AUG-1997; 97US-0906265.  
 PF  
 XX  
 XX 05-AUG-1997; 97US-0906265.  
 PR  
 XX  
 XX 01-MAY-1996; 96US-0640517.  
 PR  
 XX  
 XX 17-JUN-1996; 96US-0664487.  
 PR  
 XX  
 XX (UYJE-) UNIV JEFFERSON THOMAS.  
 PA  
 XX  
 XX Cole-Strauss AD, Kmlec EB:  
 PI  
 XX  
 XX WPI: 1999-243264/20.  
 DR  
 XX  
 XX Double-stranded oligonucleotides with containing a human wild-type  
 PT allele - useful for repairing mutations in human cells,  
 PT particularly those causing sickle cell anaemia or thalassemia  
 PS  
 XX Disclosure: Column 59-60; 40pp; English.  
 XX  
 XX The present invention describes double-stranded oligonucleotides (I)  
 CC containing fragments of wild-type human alleles. (I) are used to repair  
 CC disease associated mutations in human cells. (I) are preferably used to  
 CC treat sickle cell anaemia or thalassemia (mutations in the beta-globin  
 CC gene, including the promoter region), or Gaucher's disease (mutations in  
 CC the glucocerebrosidase gene), in haematopoietic cells. (I) may also be  
 CC used to treat familial hypercholesterolaemia (mutations in the low-  
 CC density lipoprotein receptor gene), emphysema (the alpha 1-anti-trypsin  
 CC gene), haemophilia (the factor VIII gene) or Christmas disease (the  
 CC factor IX gene), in hepatocytic cells. (I) provides repair of small  
 CC genetic mutations. The present sequence represents an oligonucleotide  
 CC from the present invention.  
 XX  
 XX Sequence 68 BP; 12 A; 18 C; 18 G; 14 T; 6 U; 0 other;  
 SQ  
 Query Match 38.8%; Score 26.4; DB 20; Length 68;  
 Best Local Similarity 65.0%; Pred. No. 2.2;  
 Matches 39; Conservative 0; Mismatches 21; Indels 0; Gaps 0;  
 QY 9 CCTACGTACGAGAAAGTTTACUUCUGCAGTACGAGUUGGAGCGCGTTTGGCGC 68  
 DB 9 CCTGAGAGAGAAAGCTGCTTTGAGUCUUCUCCCTCAGAGAGUAGUAGCGCGTTTGGCGC 68

RESULT 8  
 AAX19656  
 ID AAX19656 standard; DNA: 68 BP.  
 XX

RESULT 9  
 ABO51034  
 ID ABO51034 standard; DNA: 533 BP.  
 XX  
 XX ABO51034;  
 DE  
 XX 12-JUN-2002 (first entry)  
 DT  
 XX  
 XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 37625.  
 DE  
 XX Human; cytosine methylation; 5'-CpG-3'; unrcil; cytosine; diagnosis;  
 KW drug; side effect; cancer; central nervous system; cardiovascular;  
 KW gastrointestinal; respiratory system; single nucleotide polymorphism;  
 KW SNP; cell differentiation; ds.  
 XX

XX Homo sapiens.  
 OS  
 XX  
 PN WO200218632-A2.  
 XX  
 PD 07-MAR-2002.  
 XX  
 PF 01-SEP-2001; 2001WO-EP10074.  
 XX  
 PR 01-SEP-2000; 2000DE-1043826.  
 XX  
 PR 05-SEP-2000; 2000DE-1044543.  
 XX  
 PA (EPiG-) EPIGENOMICS AG.  
 XX  
 PI Olek A, Piepenbrock C, Berlin K, Guetig D;  
 XX  
 DR WPI: 2002-371829/40.  
 XX  
 PT Determining the degree of cytosine methylation in genomic DNA, useful  
 PT for diagnosis and prognosis, comprises selective hybridization of  
 PT amplicons from chemically treated DNA -  
 XX  
 PS Claim 12; 56pp + Sequence Listing; 56pp; German.  
 XX  
 CC This invention describes a novel method for determining the degree of  
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a  
 CC genomic sample of DNA. The sample is treated chemically to convert  
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic  
 CC DNA that contains the target C is amplified to form a labeled amplicon.  
 CC The amplicon is hybridised to two classes, each with at least one  
 CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers  
 CC and the degree of hybridisation to both classes is determined from the  
 CC label on the amplicon. From the ratio of labels hybridised to the two  
 CC classes of oligomers, the degree of methylation is calculated. The method  
 CC is used: (i) for diagnosis and/or prognosis of side effects of  
 CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders  
 CC of the central nervous, cardiovascular, gastrointestinal and respiratory  
 CC systems etc., particularly by detecting mutations or single nucleotide  
 CC polymorphisms (SNP/s); and (ii) for differentiation of cell or tissue  
 CC types and for investigating cell differentiation. The method allows the  
 CC methylation status of many C residues to be determined simultaneously.  
 CC AB013410-AB054121 represent genomic DNA sequences used to illustrate the  
 CC method for determining the degree of cytosine methylation described in  
 CC the disclosure of the invention.  
 CC  
 XX  
 SQ Sequence 533 BP; 89 A; 62 C; 208 G; 174 T; 0 other;  
 XX  
 QY Query Match 37.6%; Score 25.6; DB 24; Length 533;  
 Best Local Similarity 55.4%; Pred. No. 7.1;  
 Matches 31; Conservative 6; Mismatches 19; Indels 0; Gaps 0;  
 XX  
 Db 12 ACGTAGCAGAAAGTTTACUUCUGACGTAGGUGAAGCGCGTTTCGCG 67  
 1 ATGTCCGCGGATTTTATTATTTATGTAGCAAGGTGGAAGCCGAGATTTAAGG 56

PN WO200218632-A2.  
 XX  
 XX 07-MAR-2002.  
 PD  
 XX  
 PF 01-SEP-2001; 2001WO-EP10074.  
 XX  
 PR 01-SEP-2000; 2000DE-1043826.  
 XX  
 PR 05-SEP-2000; 2000DE-1044543.  
 XX  
 PA (EPiG-) EPIGENOMICS AG.  
 XX  
 PI Olek A, Piepenbrock C, Berlin K, Guetig D;  
 XX  
 DR WPI: 2002-371829/40.  
 XX  
 PT Determining the degree of cytosine methylation in genomic DNA, useful  
 PT for diagnosis and prognosis, comprises selective hybridization of  
 PT amplicons from chemically treated DNA -  
 XX  
 PS Claim 12; 56pp + Sequence Listing; 56pp; German.  
 XX  
 CC This invention describes a novel method for determining the degree of  
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a  
 CC genomic sample of DNA. The sample is treated chemically to convert  
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic  
 CC DNA that contains the target C is amplified to form a labeled amplicon.  
 CC The amplicon is hybridised to two classes, each with at least one  
 CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers  
 CC and the degree of hybridisation to both classes is determined from the  
 CC label on the amplicon. From the ratio of labels hybridised to the two  
 CC classes of oligomers, the degree of methylation is calculated. The method  
 CC is used: (i) for diagnosis and/or prognosis of side effects of  
 CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders  
 CC of the central nervous, cardiovascular, gastrointestinal and respiratory  
 CC systems etc., particularly by detecting mutations or single nucleotide  
 CC polymorphisms (SNP/s); and (ii) for differentiation of cell or tissue  
 CC types and for investigating cell differentiation. The method allows the  
 CC methylation status of many C residues to be determined simultaneously.  
 CC AB013410-AB054121 represent genomic DNA sequences used to illustrate the  
 CC method for determining the degree of cytosine methylation described in  
 CC the disclosure of the invention.  
 CC  
 XX  
 SQ Sequence 533 BP; 174 A; 208 C; 62 G; 89 T; 0 other;  
 XX  
 QY Query Match 37.6%; Score 25.6; DB 24; Length 533;  
 Best Local Similarity 55.4%; Pred. No. 7.1;  
 Matches 31; Conservative 6; Mismatches 19; Indels 0; Gaps 0;  
 XX  
 Db 533 ATGTCCGCGGATTTTATTATTTATGTAGCAAGGTGGAAGCCGAGATTTAAGG 478

RESULT 11  
 ABL38127  
 ID ABL38127 standard; CDNA; 435 BP.  
 XX  
 XX  
 AC ABL38127;  
 XX  
 XX  
 DT 08-APR-2002 (first entry)  
 XX  
 DE Human colon tumour antigen polynucleotide SEQ ID NO:1716.  
 XX  
 KW Human; colon cancer; colon tumour antigen; cytosolic; vaccine;  
 KW colon tumour metastatic antigen; diagnosis; gene; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200196388-A2.  
 XX  
 PD 20-DEC-2001.  
 XX  
 PF 08-JUN-2001; 2001WO-US18557.



XX 09-JUN-2000; 2000US-210899P.  
PR 20-FEB-2001; 2001US-270216P.  
XX  
PA (CORI-) CORIXA CORP.  
XX  
PI Jlang Y, Harlocker SL, Secretist H;  
XX WPI; 2002-114514/15.  
XX  
PT Novel isolated colon tumor polynucleotide differentially expressed in  
PT colon tumor or colon metastatic tumor and polypeptides encoded by them,  
XX useful for inhibiting development of cancer in patient -  
PS Claim 1; SEQ ID 1716; 105bp; English.  
XX  
CC ABI36412 to ABI3645 represent human colon tumour antigen cDNA clones (I)  
CC which were isolated from human colon tumour and colon metastatic tumour  
CC cDNA libraries. (I) have cytoskeletal activity and can be used in vaccine  
CC production. (I) can be used for stimulating and/or expanding T cells  
CC specific for a tumour protein on contact with the T cells. They are also  
CC useful for inhibiting the development of cancer in a patient. (I) can be  
CC used as probes or primers for nucleic acid hybridisation, for preparing  
CC mutant species primers, or primers for use in genetic constructions. (I)  
CC can be used in the diagnosis of a colon tumour.  
SQ  
Sequence 435 BP; 114 A; 98 C; 91 G; 130 T; 2 other;  
Query Match 37.4%; Score 25.4; DB 24; Length 435;  
Best Local Similarity 55.9%; Pred. No. 8;  
Matches 33; Conservative 5; Mismatches 21; Indels 0; Gaps 0;  
OY 2 CTTCACACCTAGCTAGCAAGTTTACUUCUGACGAGGUGAAGGCGCCT 60  
Db 104 CTTCACACCTAGCTAGCAAGTTTACUUCUGACGAGGUGAAGGCGCCT 162  
|||||  
RESULT 12  
AAI99525/C  
ID AAI99525 standard; CDNA; 1280 BP.  
XX  
AC AAI99525;  
XX  
DT 07-JAN-2002 (first entry)  
XX  
DE Human polynucleotide SEQ ID NO 23.  
XX  
KW Cytostatic; immunosuppressive; neutrotropic; neuroprotective; antiviral;  
KW antiallergic; hepatocytic; antidiabetic; antiinflammatory; antilucer;  
KW vulnerrary; anticonvulsant; antibacterial; antifungal; antiparasitic;  
KW cardiant; gene therapy; cancer; immune disorder; cardiovascular disorder;  
KW neurological disease; infection; human; secreted protein; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO200155173-A2.  
XX  
PD 02-AUG-2001.  
XX  
FE 17-JAN-2001; 2001WO-US01356.  
XX  
PR 31-JAN-2000; 2000US-0179065.  
PR 04-FEB-2000; 2000US-0180628.  
PR 24-FEB-2000; 2000US-0184664.  
PR 02-MAR-2000; 2000US-0186350.  
PR 15-MAR-2000; 2000US-0189874.  
PR 17-MAR-2000; 2000US-0190076.  
PR 18-APR-2000; 2000US-0196123.  
PR 19-MAY-2000; 2000US-0205515.  
PR 07-JUN-2000; 2000US-0209467.  
PR 28-JUN-2000; 2000US-0214886.  
PR 30-JUN-2000; 2000US-0215135.  
PR 07-JUL-2000; 2000US-0216647.  
PR

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PR 11-JUL-2000; 2000US-0217487.  
PR 11-JUL-2000; 2000US-0217486.  
PR 14-JUL-2000; 2000US-0218290.  
PR 26-JUL-2000; 2000US-0220963.  
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PR 14-AUG-2000; 2000US-0225450.  
PR 14-AUG-2000; 2000US-0225451.  
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PR 14-AUG-2000; 2000US-0225473.  
PR 14-AUG-2000; 2000US-0225474.  
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PR 14-AUG-2000; 2000US-0225480.  
PR 14-AUG-2000; 2000US-0225481.  
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PR 14-AUG-2000; 2000US-0225632.  
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PR 14-AUG-2000; 2000US-0225634.  
PR 14-AUG-2000; 2000US-0225635.  
PR 14-AUG-2000; 2000US-



CC diseases (e.g. cardiac insufficiency, coronary insufficiency or high  
CC blood pressure). The GNG DNA and protein sequences of the invention may  
CC also be used as insulin sensitizers - for improving insulin sensitivity  
CC in persons with non-insulin dependent diabetes mellitus. The present cDNA  
CC sequence encodes the human GNG-7A protein.  
XX

Sequence 2257 BP; 728 A; 522 C; 482 G; 525 T; 0 other;

Query Match 37.4%; Score 25.4; DB 24; Length 2257;  
Best Local Similarity 55.9%; Pred. No. 12;  
Matches 33; Conservative 5; Mismatches 21; Indels 0; Gaps 0;

QY 2 CTTCGACCTAGCAGAGAACTTTTACUUCGUCGAGGUGGAGGCGCGT 60  
DB 1849 CTTCGACCACTAGGAGAACTTTGATCTTGCAGCTTGATTTAATGCCACGT 1907

RESULT 14  
AAK52451

ID AAK52451 standard; cDNA; 2454 BP.

AC AAK52451;

DT 06-NOV-2001 (first entry)

DE Human polynucleotide SEQ ID NO 1980.

XX Human: cytokine; cell proliferation; cell differentiation; gene therapy;

KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;

KW tissue growth factor; immunomodulatory; cancer; leukemia;

KW nervous system disorder; arthritis; inflammation; ss.

OS Homo sapiens.

PN WO200157190-A2.

PD 09-AUG-2001.

PE 05-FEB-2001; 2001WO-US04098.

PF 03-FEB-2000; 2000US-0496914.

PR 27-APR-2000; 2000US-0560875.

PR 20-JUN-2000; 2000US-0598075.

PR 19-JUL-2000; 2000US-0620325.

PR 01-SEP-2000; 2000US-0654936.

PR 15-SEP-2000; 2000US-0663561.

PR 20-OCT-2000; 2000US-0693325.

PR 30-NOV-2000; 2000US-0728422.

XX (HYSE-) HYSEQ INC.

PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;

PI Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;

PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;

DR WPI: 2001-476283/51.

DR P-PDB: AAM79318.

XX Nucleic acids encoding polypeptides with cytokine-like activities,

XX useful in diagnosis and gene therapy -

PS Claim 1: Page 4386-4387; 6221pp; English.

XX The invention relates to polynucleotides (AAK51456-AAK53435) and the  
CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activin/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukemia, nervous system disorders, arthritis and

CC Inflammation.

CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.  
XX

Sequence 2454 BP; 782 A; 574 C; 532 G; 563 T; 3 other;

Query Match 37.4%; Score 25.4; DB 22; Length 2454;  
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QY 2 CTTCGACCTAGCAGAGAACTTTTACUUCGUCGAGGUGGAGGCGCGT 60  
DB 2175 CTTCGACCACTAGGAGAACTTTGATCTTGCAGCTTGATTTAATGCCACGT 2233

RESULT 15  
AAK52452

ID AAK52452 standard; cDNA; 2454 BP.

AC AAK52452;

DT 06-NOV-2001 (first entry)

DE Human polynucleotide SEQ ID NO 1981.

XX Human: cytokine; cell proliferation; cell differentiation; gene therapy;

KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;

KW tissue growth factor; immunomodulatory; cancer; leukemia;

KW nervous system disorder; arthritis; inflammation; ss.

OS Homo sapiens.

PN WO200157190-A2.

PD 09-AUG-2001.

PE 05-FEB-2001; 2001WO-US04098.

PF 03-FEB-2000; 2000US-0496914.

PR 27-APR-2000; 2000US-0560875.

PR 20-JUN-2000; 2000US-0598075.

PR 19-JUL-2000; 2000US-0620325.

PR 01-SEP-2000; 2000US-0654936.

PR 15-SEP-2000; 2000US-0663561.

PR 20-OCT-2000; 2000US-0693325.

PR 30-NOV-2000; 2000US-0728422.

XX (HYSE-) HYSEQ INC.

PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;

PI Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;

PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;

DR WPI: 2001-476283/51.

DR P-PDB: AAM79319.

XX Nucleic acids encoding polypeptides with cytokine-like activities,

XX useful in diagnosis and gene therapy -

PS Claim 1: Page 4387-4388; 6221pp; English.

XX The invention relates to polynucleotides (AAK51456-AAK53435) and the  
CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activin/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukemia, nervous system disorders, arthritis and

Tue Mar 18 16:16:11 2003

us-09-836-439-1.rng

Page 10

CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.

XX  
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Query Match	37.4%;	Score 25.4;	DB 22;	Length 2454;
Best Local Similarity	55.9%;	Pred. No. 15;		
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				Gaps 0;

OY 2 CTTCAACCTACGTAGACAGAAAGTTTTTACUUCUCCUACGTAGAGUGGAAGGGGGGT 60  
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 DB 2175 CTTCTGAAACACTTAGAGAAACCTTTGATCTTCAGCTTGTGATGATTTAAATGCCCACT 2233

Search completed: March 17, 2003, 10:50:28  
Job time : 393.887 secs

Gencore version 5.1.4.p5.4578  
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OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 10:23:12 ; Search time 2403.76 Seconds

(without alignments)  
458.154 Million cell updates/sec

Title: US-09-836-439-1

Perfect score: 68  
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Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Maximum Match 10%

Listing first 45 summaries

Database :

EST:\*  
1: em\_estba:\*  
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7: em\_estro:\*  
8: em\_hic:\*  
9: gp\_estl:\*  
10: gp\_estl2:\*  
11: gp\_hic:\*  
12: gp\_estl3:\*  
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25: em\_gss\_other:\*  
26: em\_gss\_pro:\*  
27: em\_gss\_rod:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

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2	27	39.7	1101	17	CNS00341
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5	25.8	37.9	899	14	B0946406 AGENCOURT
6	25.8	37.9	923	14	B0937919 AGENCOURT

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C	9	25.8	37.9	1101	12	BG295440	BG295440 602392754
C	10	25.6	37.6	288	9	AU163129	AU163129 AU163129
C	11	25.6	37.6	297	9	AU096979	AU096979 AU096979
C	12	25.6	37.6	300	10	BE401698	BE401698 CWM02E01
C	13	25.6	37.6	349	9	AU070730	AU070730 AU070730
C	14	25.6	37.6	352	9	AU057142	AU057142 AU057142
C	15	25.6	37.6	356	9	AU162789	AU162789 AU162789
C	16	25.6	37.6	358	9	AU070946	AU070946 AU070946
C	17	25.6	37.6	398	10	BE229338	BE229338 98W00005
C	18	25.6	37.6	463	9	AU057241	AU057241 AU057241
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C	20	25.6	37.6	507	10	BE230696	BE230696 99AS93 R1
C	21	25.6	37.6	573	17	AO803248	AO803248 HS_3154.A
C	22	25.6	37.6	606	17	AQ049317	AQ049317 CLM-13E4.A
C	23	25.6	37.6	909	9	AL566335	AL566335 AL566335
C	24	25.4	37.4	432	12	BF149104	BF149104 35_11 Hum
C	25	25.4	37.4	464	10	AW377397	AW377397 QVO-CT022
C	26	25.4	37.4	466	9	AA625142	AA625142 aE70C02.r
C	27	25.4	37.4	472	12	BF048056	BF048056 dc88H10.Y
C	28	25.4	37.4	494	14	BM706039	BM706039 UT-E-DW0-
C	29	25.4	37.4	510	10	AW022120	AW022120 dT33b07.Y
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C	31	25.4	37.4	576	14	BM696870	BM696870 UT-E-DW0-
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C	38	25.4	37.4	675	13	B1596522	B1596522 603243571
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## ALIGNMENTS

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DEFINITION RST7364 Athersys RAGE Library Homo sapiens cDNA, mRNA sequence.  
ACCESSION BG188346  
VERSION BG188346.1 GI:13710033  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 818)  
Harrington,J.J., Sherf,B., Rundlett,S., Jackson,P.D., Perry,R., Cain,S., Leventhal,C., Thornton,M., Ramchandran,R., Whittington,J., Lerner,U., Coslano,D., McElligott,K., Booser,S., Maye,R., Smith,E., Veloso,N., Kika,A., Hess,J., Cothren,K., Lo,K., Offenbacher,J., Danzig,J., and Ducar,M.  
Creation of genome-wide protein expression libraries using random activation of gene expression  
Nat. Biotechnol. 19 (5), 440-445 (2001)  
21227151  
JOURNAL MEDLINE  
COMMENT Contact: Scott J. Cain  
Athersys, Inc.  
3201 Carnegie Ave, Cleveland, OH 44115, USA  
Tel: 216 431 9900  
Fax: 216 361 9596  
Email: scai@atersys.com  
High quality sequence stop: 440.  
Location/Qualifiers

	source	1..818 /organism="Homo sapiens" /db_xref="taxon:9606" /clone_lib="Athersys RAGE Library" /cell_line="HT1080" /note="See 'Creation of Genome-wide Protein Expression Libraries using Random Activation of Gene Expression', Nature Biotechnology, in press. Note that even though the cell type indicated is HT1080, since a random activation method was used, these sequence tags are not necessarily expressed in HT1080 under normal circumstances."
BASE COUNT	218 a 197 c 181 g 222 t	
ORIGIN		
Query Match	39.7% Best Local Similarity 57.6% Matches 34; Conservative 5;	Score 27; DB 12; Length 818; Pred. No. 17; Mismatches 20; Indels 0; Gaps 0;
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RESULT 2	CNS0034I	1101 bp DNA linear GSS 03-JUN-1999
LOCUS	Drosophila melanogaster genome survey sequence TET3 end of BAC # BACR07N08 of RPCI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.	
ACCSSION	AL063844	
VERSION	AL063844.1 GI:4941600	
KEYWORDS	GSS.	
SOURCE	Drosophila melanogaster.	
ORGANISM	Drosophila melanogaster. Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.	
REFERENCE	1 (bases 1 to 1101) Genoscope.	
AUTHORS	Direct Submission	
TITLE	Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage ; BP 191 91006 EVRY cedex - FRANCE (E-mail : sequef@genoscope.cns.fr	
JOURNAL	- Web : www.genoscope.cns.fr)	
COMMENT	Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see http://www.fruitfly.org The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osoegawa and Aaron Mammoser in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of drosophila DNA provided by the BDGP from the isogenic strain y2; cn bw sp. the same strain used for the BDGP's pl and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.	
FEATURES	Location/Qualifiers	
source	1..1101 /organism="Drosophila melanogaster" /db_xref="taxon:7227" /clone="BACR07N08" /clone_lib="RPCI-98" /note="end : TET3"	
BASE COUNT	349 a 179 c 216 g 325 t 32 others	
ORIGIN		
Query Match	39.7% Best Local Similarity 58.8% Matches 30; Conservative 6;	Score 27; DB 17; Length 1101; Pred. No. 19; Mismatches 15; Indels 0; Gaps 0;
Y	15 TAGCAGAAAGTTTTTACUUCUGCUCACGTAGGUAAGGCGCGTTTTCG 65	

DB	693	TTCGATTAAGTTTCTTACTGTTACTGATGATTTTGGAGTGGGTTTGTGG	743
RESULT 3			
LOCUS	B1739136/c		
DEFINITION	B1739136	741 bp	mRNA
ACCESSION	603361404F1 NIH_MGC_94	Mus musculus	cdna clone IMAGE:5368368 5',
VERSION	B1739136		
KEYWORDS	B1739136.1	GI:15716149	
SOURCE	EST.		
ORGANISM	house mouse.		
	Mus musculus		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.		
TITLE	1 (bases 1 to 741)		
JOURNAL	NIH-MGC <a href="http://mgc.nci.nih.gov/">http://mgc.nci.nih.gov/</a> .		
COMMENT	National Institutes of Health, Mammalian Gene Collection (MGC)		
	Unpublished (1999)		
	Contact: Robert Strausberg, Ph.D.		
	Email: <a href="mailto:cgabs-remail.nih.gov">cgabs-remail.nih.gov</a>		
	Tissue Procurement: The Cepko Laboratory		
	cDNA Library Preparation: Life Technologies, Inc.		
	cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)		
	DNA Sequencing by: Incyte Genomics, Inc.		
	Clone distribution: MGC clone distribution information can be		
	found through the I.M.A.G.E. Consortium/LNL at:		
	<a href="http://image.lnl.gov">http://image.lnl.gov</a>		
	Plate: L1LM11937 row: h column: 01		
	High quality sequence stop: 729.		
FEATURES	Location/Qualifiers		
Source	1..741		
	/organism="Mus musculus"		
	/db_xref="taxon:10090"		
	/clone="IMAGE:5368368"		
	/clone_lib="NIH_MGC_94"		
	/tissue_type="retina"		
	/lab_host="DH10B (phage-resistant)"		
	/note="Organ: eye; Vector: pCMV-Sport6; Site_1: NotI;		
	Site_2: SalI; Cloned unidirectionally; oligo-dT primed.		
	Average insert size 3.3 kb. Library enriched for		
	full-length clones and constructed by Life Technologies.		
	Note: this is a NIH-MGC Library."		
BASE COUNT	210 a	177 c	194 g
ORIGIN		160 t	
Query Match		37.9%	Score 25.8; DB 13; Length 741;
Best Local Similarity		69.0%;	Pred. No. 47;
Matches 20; Conservative		7; Mismatches	2; Indels 0; Gaps 0;
QY	30	ACUUUCUGCUACGACGAGUGGAGGCGCC	58
	:     :     :		
Db	368	ACTTCTGCTACGATGAGTGGAGGCCAC	340
RESULT 4			
LOCUS	B1733738/c	747 bp	mRNA
DEFINITION	60335355F1 NIH_MGC_94	Mus musculus	cdna clone IMAGE:5359908 5',
ACCESSION	B1733738		
VERSION	B1733738		
KEYWORDS	B1733738.1	GI:15710751	
SOURCE	EST.		
ORGANISM	house mouse.		
	Mus musculus		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.		
TITLE	1 (bases 1 to 747)		
JOURNAL	NIH-MGC <a href="http://mgc.nci.nih.gov/">http://mgc.nci.nih.gov/</a> .		
COMMENT	National Institutes of Health, Mammalian Gene Collection (MGC)		
	Unpublished (1999)		
	Contact: Robert Strausberg, Ph.D.		

Email: c9apbs-remail.nih.gov  
Tissue Procurement: The Cepko Laboratory  
cDNA Library Preparation: Life Technologies, Inc.  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:  
http://image.llnl.gov  
Plate: LLM13915 row: g column: 13  
High quality sequence stop: 734.

## FEATURES

Location/Qualifiers  
1..747

/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:5359908"  
/clone\_lib="NIH\_MGC\_94"  
/tissue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: pCMV-SPORT6; Site:1: NotI;  
Site:2: SalI; Cloned unidirectionally; oligo-dT primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."

BASE COUNT  
218 a 169 c 198 g 162 t

## ORIGIN

## Query Match

Best Local Similarity 69.0%; Pred. No. 47;  
Matches 20; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

Qy 30 ACUUCUCGACGTAGGUGGAGGCGC 58

Db 183 ACTTCTGCTACGTAGGTGGAAGCCAC 155

## RESULT 5

BQ946406 899 bp mRNA linear EST 21-AUG-2002

## LOCUS

BQ946406 AGENCOURT\_8924053 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:6467512

## DEFINITION

5', mRNA sequence.

## ACCESSION

BQ946406

## VERSION

BQ946406.1

## KEYWORDS

GI:22361884

## SOURCE

EST.

## ORGANISM

house mouse.

## REFERENCE

1 (bases 1 to 899)

## AUTHORS

NIH-MGC http://mgc.ncl.nih.gov/.

## TITLE

Unpublished (1999)

## JOURNAL

Contact: Robert Strausberg, Ph.D.

## COMMENT

Email: c9apbs-remail.nih.gov

## Tissue Procurement

The Cepko Laboratory

## cDNA Library Preparation

Life Technologies, Inc.

## DNA Sequencing by

The I.M.A.G.E. Consortium (LNLN)

## Clone distribution

MGC clone distribution information can be

## found through the I.M.A.G.E. Consortium/LNLN at:

http://image.llnl.gov

## Plate

LLM13915 row: 1 column: 17

## High quality sequence stop

614.

## FEATURES

Location/Qualifiers  
1..899

/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:6467512"  
/clone\_lib="NIH\_MGC\_94"  
/tissue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: pCMV-SPORT6; Site:1: NotI;  
Site:2: SalI; Cloned unidirectionally; oligo-dT primed.  
Average insert size 3.3 kb. Library enriched for

Full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."

BASE COUNT  
248 a 208 c 231 g 207 t 5 others

## ORIGIN

## Query Match

Best Local Similarity 69.0%; Pred. No. 50;  
Matches 20; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

Qy 30 ACUUCUCGACGTAGGUGGAGGCGC 58

Db 532 ACTTCTGCTACGTAGGTGGAAGCCAC 504

## RESULT 6

BQ937919 923 bp mRNA linear EST 21-AUG-2002

## LOCUS

BQ937919 AGENCOURT\_8932027 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:6465857

## DEFINITION

5', mRNA sequence.

## ACCESSION

BQ937919

## VERSION

BQ937919.1

## KEYWORDS

GI:22353397

## SOURCE

EST.

## ORGANISM

house mouse.

## REFERENCE

1 (bases 1 to 923)

## AUTHORS

NIH-MGC http://mgc.ncl.nih.gov/.

## TITLE

Unpublished (1999)

## JOURNAL

Contact: Robert Strausberg, Ph.D.

## COMMENT

Email: c9apbs-remail.nih.gov

## Tissue Procurement

The Cepko Laboratory

## cDNA Library Preparation

Life Technologies, Inc.

## DNA Sequencing by

The I.M.A.G.E. Consortium (LNLN)

## Clone distribution

MGC clone distribution information can be

## found through the I.M.A.G.E. Consortium/LNLN at:

http://image.llnl.gov

## Plate

LLM13989 row: d column: 18

## High quality sequence stop

666.

## FEATURES

Location/Qualifiers  
1..923

## source

/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:6465857"  
/clone\_lib="NIH\_MGC\_94"  
/tissue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: pCMV-SPORT6; Site:1: NotI;  
Site:2: SalI; Cloned unidirectionally; oligo-dT primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."

## BASE COUNT

251 a 224 c 233 g 214 t 1 others

## ORIGIN

Query Match

## Best Local Similarity

69.0%; Pred. No. 50;

## Matches

20; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

Qy 30 ACUUCUCGACGTAGGUGGAGGCGC 58

Db 667 ACTTCTGCTACGTAGGTGGAAGCCAC 639

## RESULT 7

BG262012 943 bp mRNA linear EST 13-FEB-2001

## LOCUS

BG262012 60233784F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:4481291 5'

## DEFINITION

mRNA sequence.

## ACCESSION

BG262012

## VERSION

BG262012.1

GI:12771828

KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
1 (bases 1 to 943)  
REFERENCE NIH-MGC <http://mhc.nci.nih.gov/>.  
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)  
TITLE Unpublished (1999)  
JOURNAL Contact: Robert Strausberg, Ph.D.  
COMMENT Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
Tissue Procurement: The Cepko Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>  
Plate: LLAM10316 row: f column: 12  
High quality sequence stop: 694.  
Location/Qualifiers  
1. .943  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:4481943"  
/clone\_lib="NIH\_MGC\_94"  
/tissue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: pCMV-SPORE6; Site\_1: NotI;  
Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."

BASE COUNT 269 a 230 c 256 g 188 t  
ORIGIN

Query Match 37.9%; Score 25.6; DB 12; Length 943;  
Best Local Similarity 69.0%; Pred. No. 51;  
Matches 20; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

QY 30 ACUUCUGCUCAGTAGGUGAAGGCCGC 58  
||||:||||:||||:||||:||||:|  
Db 471 ACTTCTGCTAGCTAGGTGGAGGCCAC 443

RESULT 8  
BG342737/c 961 bp mRNA linear EST 27-FEB-2001  
DEFINITION 602374557F1 NIH\_MGC\_94 Mus musculus CDNA clone IMAGE:4481943 5',  
LOCUS mRNA sequence.  
ACCESSION BG342737  
VERSION BG342737.1 GI:13149175  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 961)  
REFERENCE NIH-MGC <http://mhc.nci.nih.gov/>.  
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)  
TITLE Unpublished (1999)  
JOURNAL Contact: Robert Strausberg, Ph.D.  
COMMENT Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
Tissue Procurement: The Cepko Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>  
Plate: LLAM10318 row: a column: 16  
High quality sequence stop: 653.  
Location/Qualifiers

FEATURES

source 1. .961  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:4481943"  
/clone\_lib="NIH\_MGC\_94"  
/tissue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: pCMV-SPORE6; Site\_1: NotI;  
Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."

BASE COUNT 288 a 218 c 254 g 201 t  
ORIGIN

Query Match 37.9%; Score 25.8; DB 12; Length 961;  
Best Local Similarity 69.0%; Pred. No. 51;  
Matches 20; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

QY 30 ACUUCUGCUCAGTAGGUGAAGGCCGC 58  
||||:||||:||||:||||:||||:|  
Db 31 ACTTCTGCTAGCTAGGTGGAGGCCAC 3

RESULT 9  
BG295440/c 1101 bp mRNA linear EST 21-FEB-2001  
DEFINITION 602392754F1 NIH\_MGC\_94 Mus musculus CDNA clone IMAGE:4504476 5',  
LOCUS mRNA sequence.  
ACCESSION BG295440  
VERSION BG295440.1 GI:13057077  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 1101)  
REFERENCE NIH-MGC <http://mhc.nci.nih.gov/>.  
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)  
TITLE Unpublished (1999)  
JOURNAL Contact: Robert Strausberg, Ph.D.  
COMMENT Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
Tissue Procurement: The Cepko Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>  
Plate: LLAM10376 row: 1 column: 13  
High quality sequence stop: 735.  
Location/Qualifiers  
1. .1101  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:4504476"  
/clone\_lib="NIH\_MGC\_94"  
/tissue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: pCMV-SPORE6; Site\_1: NotI;  
Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."

BASE COUNT 329 a 279 c 289 g 204 t  
ORIGIN

Query Match 37.9%; Score 25.8; DB 12; Length 1101;  
Best Local Similarity 69.0%; Pred. No. 54;  
Matches 20; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

QY 30 ACUUCUGCUCAGTAGGUGAAGGCCGC 58  
||||:||||:||||:||||:||||:|



REFERENCE	1 (bases 1 to 288)
AUTHORS	Sasaki, T. and Yamamoto, K.
TITLE	Rice cDNA from green shoot (2000)
JOURNAL	Unpublished (2000)
COMMENT	Contact: Takuji Sasaki

```
PROJECT = 'RGP', url=http://rgp.dna.afric.go.jp/
FEATURES
  source      Location/Qualifiers
1. 288
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BASE COUNT	ORIGIN	83 a	79 c	68 g	57 t	1 others
"/notes"Green shoot (8 days old)"						
"/along"rice green shoot"						

Query Match	37.6%	Score 25.6;	DB 9;	Length 288;
Best Local Similarity	60.4%;	Pred. No. 41;		
Matches	29; Conservative	5; Mismatches	14;	Indels

QY 8 ACCTAGTACGACGAAAGTTTTCACUUCUCGACCTAGUGGAGGG 55  
 || ||||| || ||| ||| :: : ||||| | - ||  
 Db 210 ACCTAGTACGTACCGACGATTTATTAGTAGTACCTAGTACGATGG 163

RESULT 11	
AU096979/c	
LOCUS	AU096979
DEFINITION	297 bp mRNA linear EST 03-APR-2007
CDNA clone S16784,	Rice green shoot Oryza sativa (japonica cultivar-group)
VERSION	AU096979
ACCESSION	AU096979.1 GI:8859661
KEYWORDS	EST.
SOURCE	Oryza sativa (japonica cultivar-group).
ORGANISM	Oryza sativa (japonica cultivar-group).

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
1 (bases 1 to 297)	Sasaki, T. and Yamamoto, K.	Rice cDNA from green shoot (2000)	Unpublished (2000)	
Contact:	Takui Sasaki			
	National Institute of Agrobiological Resources			
	Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki			
	305-8602, Japan			
	Tel.: 81-298-38-7441			
	Fax: 81-298-38-7468			
	Email: tsasaki@agr.affrc.go.jp, URL: <a href="http://rgp.dna.affrc.go.jp/">http://rgp.dna.affrc.go.jp/</a>			
	PROJECT "RGP."			

FEATURES	Location/Qualifiers
source	1. .297

BASE COUNT	ORIGIN	clone_11b="Rice green shoot" /note="Green shoot (8 days old)"
83	a	84 c
		72 g
		56 t
		2 others

Query Match	37.6%	Score 25.6	DB 9	Length 297
Best Local Similarity	60.4%	Pred. No 41		
Matches 29: Conservative	5	Mismatches 14	Indels 0	Gaps 0
QY	8	ACCTACGTAGCAGAAAGTTTACUUCUUCAGCTAGAGUGGAGGCG	55	
Db	219	ACGTACGTAGTACCGACGATTATTATTAGTGGTAGCTAGTCGATGCG	172	

RESULT 12	
BE401698/c	
LOCUS	300 bp mRNA linear EST 21-JUL-2000
DEFINITION	aestivum CDNA clone CWM02EL010, mRNA sequence.
ACCESSION	BE401698
VERSION	BE401698.1 GI:9361166
KEYWORDS	EST.
SOURCE	bread wheat.

ORGANISM  
Triticum aestivum  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae  
; Triticeae; Triticum.  
1 (bases 1 to 300)  
REFERENCE  
Anderson, C.A., Appels, R., Bailey, P., Blake, T., Close, T., Cloutier

TITLE  
International Trillaceae EST Cooperative (INEC): Production of  
Expressed Sequence Tags for Species of the Trillaceae  
Unpublished (2000)  
JOURNAL  
Contact: jia\_j  
COMMENT

key lab. of Crop Germplasm & Biotechnology  
Inst. of Crop Germplasm Resources  
Beijing 100081 PR CHINA  
Tel: 86 1 62186623  
Fax: 86 1 62186629  
Email: jileng@iml.cnc.ac.cn  
International Triflicae EST Cooperative (ITEC)  
[http://whmat.pw.usda.gov/genome.  
location/duan/jif/](http://whmat.pw.usda.gov/genome.location/duan/jif/)

FEATURES	
Location/Qualifiers	1. .300

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/organism="Triticum aestivum"
/cultivar="Powdery Mildew Resistant line"
/db_xref="taxon:4565"
/clone="CMM02EL010"
/clone_1lb="ITEC CMW Wheat Powdery Mildew Resistant
Library"
/tissue_type="leaf"
/dev_stage="seedling, challenged with powdery mildew
strain"
/notes="Vector: Lambda Triplex, site 1: SfiI, primer used
5'-TCCGACATCTGACGAGC-3', 500 bp average insert size."
BASE COUNT      84 a
                81 c      87 g      48 t
                81 c      87 g      48 t

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Query Match	37.6%	Score 25.6	DB 10	length 300
Best Local Similarity	60.4%	Pred. No. 41		
Matches 29	Conservative 5	Mismatches 14	Indels 0	Gaps 0
8 ACCTAGTAGCAGAAAGTTTACUUCUCUACGTAAGUUGAAGCG 55				

Db 260 ACGTACGTAGCAGCATATTATTAGTGTAGTACGTACGTACGATGG 213

RESULT 13  
A0070730 349 bp mRNA linear EST 02-APR-2002

LOCUS A0070730 Rice cDNA from young root Oryza sativa (japonica  
DEFINITION cultivar-group) cDNA clone R10161\_2A, mRNA sequence.

ACCESSION A0070730.1 GI:5038620

VERSION EST.  
KEYWORDS Oryza sativa (japonica cultivar-group).  
SOURCE Oryza sativa (japonica cultivar-group).  
ORGANISM Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
Ehrhartoideae; Oryzaceae; Oryza.

REFERENCE 1 (bases 1 to 349)  
AUTHORS Yamamoto, K. and Sasaki, T.  
TITLE Rice cDNA from young root  
JOURNAL Unpublished (1999)  
COMMENT Contact: Takuji Sasaki  
National Institute of Agrobiological Resources  
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki  
305-8602, Japan  
Tel: 81-298-38-7441  
Fax: 81-298-38-7468  
Email: tsasaki@abr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/  
PROJECT = "RGP"

FEATURES  
source Location/Qualifiers

1..349  
/organism="Oryza sativa (japonica cultivar-group)"  
/cultivar="Nipponbare"  
/db\_xref="taxon:39947"  
/clone="R10161\_2A"  
/clone\_1ib="Rice cDNA from young root"  
/tissue\_type="young root"  
BASE COUNT 58 a 98 c 96 g 97 t  
ORIGIN

Query Match 37.6%; Score 25.6; DB 9; Length 349;  
Best Local Similarity 60.4%; Pred. No. 43;  
Matches 29; Conservative 5; Mismatches 14; Indels 0; Gaps 0;

OY 8 ACGTACGTAGCAGCAAGTTTACUUCUGCUCAGTACGTAGGAGG 55  
Db 85 ACGTACGTAGCAGCATATTATTAGTGTAGTACGTACGATGG 132

RESULT 14  
A0057142/c 352 bp mRNA linear EST 01-APR-2002  
LOCUS A0057142 Oryza sativa mature leaf Nipponbare Oryza sativa (japonica  
DEFINITION cultivar-group) cDNA clone S21188\_1A, mRNA sequence.

ACCESSION A0057142.1 GI:4716026  
VERSION EST.  
KEYWORDS Oryza sativa (japonica cultivar-group).  
SOURCE Oryza sativa (japonica cultivar-group).  
ORGANISM Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
Ehrhartoideae; Oryzaceae; Oryza.  
1 (bases 1 to 352)  
Yamamoto, K. and Sasaki, T.  
Rice cDNA from mature leaf  
Unpublished (1999)  
Contact: Takuji Sasaki  
National Institute of Agrobiological Resources  
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki  
305-8602, Japan  
Tel: 81-298-38-7441  
Fax: 81-298-38-7468  
Email: tsasaki@abr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/

PROJECT = "RGP"  
FEATURES Location/Qualifiers  
source 1..352  
/organism="Oryza sativa (japonica cultivar-group)"  
/cultivar="Nipponbare"  
/db\_xref="taxon:39947"  
/clone="S21188\_1A"  
/clone\_1ib="Oryza sativa mature leaf Nipponbare"  
/tissue\_type="mature leaf"

BASE COUNT 97 a 97 c 100 g 58 t  
ORIGIN

Query Match 37.6%; Score 25.6; DB 9; Length 352;  
Best Local Similarity 60.4%; Pred. No. 44;  
Matches 29; Conservative 5; Mismatches 14; Indels 0; Gaps 0;

OY 8 ACGTACGTAGCAGCAAGTTTACUUCUGCUCAGTACGTAGGAGG 55  
Db 268 ACGTACGTAGCAGCATATTATTAGTGTAGTACGTACGATGG 221

RESULT 15  
A0162789/c 356 bp mRNA linear EST 03-APR-2002  
LOCUS A0162789 Rice mature leaf Oryza sativa (japonica cultivar-group)  
DEFINITION cDNA clone S21904, mRNA sequence.

ACCESSION A0162789.1 GI:11026188  
VERSION EST.  
KEYWORDS Oryza sativa (japonica cultivar-group).  
SOURCE Oryza sativa (japonica cultivar-group).  
ORGANISM Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
Ehrhartoideae; Oryzaceae; Oryza.

REFERENCE 1 (bases 1 to 356)  
AUTHORS Sasaki, T. and Yamamoto, K.  
TITLE Rice cDNA from mature leaf (2000)  
JOURNAL Unpublished (2000)  
COMMENT Contact: Takuji Sasaki  
National Institute of Agrobiological Resources  
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki  
305-8602, Japan  
Tel: 81-298-38-7441  
Fax: 81-298-38-7468  
Email: tsasaki@abr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/  
PROJECT = "RGP"

FEATURES Location/Qualifiers  
source 1..356  
/organism="Oryza sativa (japonica cultivar-group)"  
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BASE COUNT 98 a 99 c 99 g 58 t 2 others  
ORIGIN

Query Match 37.6%; Score 25.6; DB 9; Length 356;  
Best Local Similarity 60.4%; Pred. No. 44;  
Matches 29; Conservative 5; Mismatches 14; Indels 0; Gaps 0;

OY 8 ACGTACGTAGCAGCAAGTTTACUUCUGCUCAGTACGTAGGAGG 55  
Db 272 ACGTACGTAGCAGCATATTATTAGTGTAGTACGTACGATGG 225

Search completed: March 17, 2003, 13:09:07  
Job time: 2409.76 secs

GenCore version 5.1.4-p5\_4578  
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# OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 09:43:42 ; Search time 230.108 Seconds  
(without alignments)  
3161.870 Million cell updates/sec

Title: US-09-836-439-2

Perfect score: 25  
Sequence: 1 ccatgacacattggaatgagag 25

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

## Database :

GenEmbl:\*  
1: gb.ba:\*  
2: gb.htg:\*  
3: gb.in:\*  
4: gb.om:\*  
5: gb.ov:\*  
6: gb.pat:\*  
7: gb.ph:\*  
8: gb.pl:\*  
9: gb.pr:\*  
10: gb.ro:\*  
11: gb.sts:\*  
12: gb.sy:\*  
13: gb.un:\*  
14: gb.vl:\*  
15: em.ba:\*  
16: em.fun:\*  
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18: em.in:\*  
19: em.mu:\*  
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34: em.htg.pln:\*  
35: em.htg.rod:\*  
36: em.htg.man:\*  
37: em.htg.vtc:\*  
38: em.sy:\*  
39: em.htgo.hum:\*  
40: em.htgo.mus:\*  
41: em.htgo.other:\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	23.4	93.6	85	10	MMHIFLAS04
2	23.4	93.6	538	6	AF004144 Mus muscu
3	23.4	93.6	2481	6	AF004144 Mus muscu
4	23.4	93.6	2481	6	AF004144 Mus muscu
5	23.4	93.6	2509	6	AF004144 Mus muscu
6	23.4	93.6	2522	6	AF004144 Mus muscu
7	23.4	93.6	2528	6	AF004144 Mus muscu
8	23.4	93.6	2530	6	AF004144 Mus muscu
9	23.4	93.6	2537	6	AF004144 Mus muscu
10	23.4	93.6	2711	10	MMHIFLAS04
11	23.4	93.6	3551	9	AF004144 Mus muscu
12	23.4	93.6	3678	9	AF004144 Mus muscu
13	23.4	93.6	3718	10	AF004144 Mus muscu
14	23.4	93.6	3745	6	AF004144 Mus muscu
15	23.4	93.6	3745	10	MMHIFLAS04
16	23.4	93.6	3867	10	MMHIFLAS04
17	23.4	93.6	3927	6	AF004144 Mus muscu
18	23.4	93.6	3933	9	AF004144 Mus muscu
19	23.4	93.6	3942	9	AF004144 Mus muscu
20	23.4	93.6	3973	10	AF004144 Mus muscu
21	23.4	93.6	4183	10	AF004144 Mus muscu
22	23.4	93.6	10355	6	AF004144 Mus muscu
23	23.4	93.6	20463	10	MMHIFLAS04
24	23.4	93.6	169792	2	AC104313
25	23.4	93.6	188107	2	CNSOLDME
26	23.4	93.6	228786	2	AC124712
27	21.8	87.2	2551	4	AB018398
28	20.4	81.6	171838	2	AC105321
29	20.2	80.8	3006	5	AF02782
30	20.2	80.8	3076	5	AF212989
31	20.2	80.8	117771	9	AC112906
32	20.2	80.8	181825	2	AC068191
33	19.8	79.2	141762	2	HS198C21
34	19.8	79.2	14837	2	AC021509
35	19.8	79.2	165840	9	AC068112
36	19.8	79.2	187277	9	AC022968
37	19.8	79.2	200000	2	AP000494
38	19.8	79.2	200567	2	AP001587
39	19.8	79.2	203540	2	AP002396
40	19.4	77.6	1596	8	VSCL308P
41	19.4	77.6	2110	8	SCYGR148C
42	19.4	77.6	2679	8	SCYGR149W
43	19.4	77.6	2677	8	SCYGR149W
44	19.4	77.6	201802	2	AC124473
45	19.4	77.6	264009	2	AC115294

## ALIGNMENTS

RESULT 1  
LOCUS MMHIFLAS04  
DEFINITION Mus musculus hypoxia-inducible factor 1 alpha (Hif1a) gene, exon 4.  
ACCESSION AF004144  
VERSION AF004144.1 GI:2197137  
KEYWORDS  
SEGMENT  
SOURCE  
ORGANISM  
Mus musculus.  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE  
Luo, G., Gu, Y. Z., Jain, S., Chan, W. K., Carr, K. M., Hogenesch, J. B. and  
Bradfield, C. A.

TITLE Molecular characterization of the murine Hif-1 alpha locus  
JOURNAL Gene Expr. 6 (5), 287-299 (1997)  
MEDLINE 98034461  
PUBMED 9368100  
REFERENCE 2 (bases 1 to 85)  
AUTHORS Luo, G., Gu, X., -Z., Jain, S., Chan, W.K., Carr, K.M., Hogenesch, J.B. and Bradfield, C.A.  
JOURNAL Submitted (14-MAY-1997) Oncology, University of Wisconsin-Madison, 1400 University Ave., Madison, WI 53706, USA  
FEATURES  
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Best Local Similarity 96.0%; Pred. No. 0.43;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
OY 1 CCATGTGACCATGAGGAATGAGAG 25  
Db 40 CCATGTGACCATGAGGAATGAGAG 64  
RESULT 2  
LOCUS HSHF1A03 538 bp DNA linear PRI 26-OCT-1998  
DEFINITION Homo sapiens hypoxia-inducible factor 1 alpha subunit (HIF1A) gene,  
exons 3 and 4.  
ACCESSION AF050117  
VERSION AF050117.1 GI:3790523  
KEYWORDS 3 of 13  
SEGMENT  
SOURCE Homo sapiens.  
ORGANISM Homo sapiens.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 538)  
AUTHORS Iyer, N.V., Leung, S.W. and Semenza, G.L.  
TITLE The human hypoxia-inducible factor 1alpha gene: HIF1A structure and evolutionary conservation  
JOURNAL Genomics 52 (2), 159-165 (1998)  
MEDLINE 99000835  
PUBMED 9782081  
REFERENCE 2 (bases 1 to 538)  
AUTHORS Iyer, N.V., Leung, S.W. and Semenza, G.L.  
TITLE Direct Submission  
JOURNAL Submitted (24-FEB-1998) Departments of Pediatrics and Medicine, Institute of Genetic Medicine, Johns Hopkins University School of Medicine, 600 N. Wolfe St, Baltimore, MD 21287-3914, USA  
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OY 1 CCATGTGACCATGAGGAATGAGAG 25  
Db 337 CCATGTGACCATGAGGAATGAGAG 361  
RESULT 3  
LOCUS AX451938 2481 bp DNA linear PAT 03-JUL-2002  
DEFINITION Sequence 3 from Patent WO0212326.  
ACCESSION AX451938  
VERSION AX451938.1 GI:21698761  
KEYWORDS human.  
SOURCE  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1  
AUTHORS Poellinger, L., Pereira, T. and Ruas, J.  
TITLE Mechanism of conditional regulation of the hypoxia-inducible factor-1 by the von hippel-lindau tumor suppressor protein  
JOURNAL Patent: WO 0212326-A 3 14-FEB-2002;  
Angiogenetics Sweden AB (SE)  
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Best Local Similarity 96.0%; Pred. No. 0.38;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
OY 1 CCATGTGACCATGAGGAATGAGAG 25  
Db 412 CCATGTGACCATGAGGAATGAGAG 436  
RESULT 4  
LOCUS AX481424 2481 bp DNA linear PAT 16-AUG-2002  
DEFINITION Sequence 38 from Patent WO02055693.  
ACCESSION AX481424

VERSION AX481424.1 GI:22316338  
 KEYWORDS human.  
 SOURCE Homo sapiens  
 ORGANISM Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

REFERENCE 1  
 AUTHORS Kreutzer, R., Limmer, S., Rost, S. and Hadwiger, P.  
 TITLE Method for inhibiting the expression of a target gene  
 JOURNAL Patient: WO 02055693-A 38 18-JUL-2002;  
 Ribopharma AG (DE)  
 FEATURES  
 Source Location/Qualifiers  
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BASE COUNT 829 a 512 c 500 g 640 t  
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Query Match 93.6%; Score 23.4; DB 6; Length 2481;  
 Best Local Similarity 96.0%; Pred. No. 0.38;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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 DB 412 CCATGTGACCATGGAATGAGAG 436

RESULT 5  
 AF304431 2509 bp mRNA linear PRI 29-DEC-2000  
 LOCUS Homo sapiens hypoxia-inducible factor 1 alpha subunit (HIF1A) mRNA,  
 DEFINITION complete cds.  
 ACCESSION AF304431  
 VERSION AF304431.1 GI:11995454  
 KEYWORDS  
 SOURCE  
 ORGANISM Homo sapiens.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

REFERENCE 1 (bases 1 to 2509)  
 AUTHORS Sun, B., Zhao, H.R., Yu, R.T. and Ni, M.S.H.  
 TITLE Direct Submission  
 JOURNAL Submitted (11-SEP-2000) Department of Neurosurgery, Affiliated  
 Hospital of Xuzhou Medical College, Huaihai West Road, Xuzhou,  
 Jiangsu 221002, China  
 FEATURES  
 Source Location/Qualifiers  
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 ORIGIN

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 Best Local Similarity 96.0%; Pred. No. 0.38;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCATGTGACCATGGAATGAGAG 25  
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 DB 412 CCATGTGACCATGGAATGAGAG 436

RESULT 6  
 AF208487 2522 bp DNA linear PRI 27-DEC-1999  
 LOCUS Homo sapiens hypoxia-inducible factor 1 alpha (HIF1A) gene,  
 DEFINITION complete cds.  
 ACCESSION AF208487  
 VERSION AF208487.1 GI:6636337  
 KEYWORDS  
 SOURCE  
 ORGANISM Homo sapiens.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

REFERENCE 1 (bases 1 to 2522)  
 AUTHORS Rupert, J.L. and Hochachka, P.W.  
 TITLE HIF1A sequence in the Quechua, a high altitude population  
 JOURNAL Unpublished  
 2 (bases 1 to 2522)  
 REFERENCE Rupert, J.L. and Hochachka, P.W.  
 TITLE Direct Submission  
 JOURNAL Submitted (25-NOV-1999) Zoology, University of British Columbia,  
 6270 University Blvd., Vancouver, BC V6T 1Z4, Canada  
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QY 1	CCATGTGACCATTTAGGAATGAGAG 25				
Db 535	CCATGTGACCATTTAGGAATGAGAG 559				
RESULT 11					
LOCUS	AB073325				
DEFINITION	Homo sapiens HIF1A mRNA for hypoxia-inducible factor 1 alpha				
ACCESSION	AB073325				
VERSION	AB073325.1				
KEYWORDS					
SOURCE	Homo sapiens liver cDNA to mRNA.				
ORGANISM	Homo sapiens				
REFERENCE					
AUTHORS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
TITLE	Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.				
JOURNAL					
REFERENCE					
AUTHORS	Tanaka,S. and Sugimachi,K.				
TITLE	Hypoxia-inducible factor-1 alpha variant isolated from human liver				
JOURNAL					
REFERENCE					
AUTHORS	Unpublished				
TITLE	2 (bases 1 to 3551)				
JOURNAL					
REFERENCE					
AUTHORS	Tanaka,S. and Sugimachi,K.				
TITLE	Direct Submission				
JOURNAL					
REFERENCE					
AUTHORS	Submitted (20-OCT-2001) Shinji Tanaka, Kyushu University, Graduate				
TITLE	School of Medical Sciences, Department of Surgery and Science;				
JOURNAL					
REFERENCE					
AUTHORS	3-1-1 Maizashi, Fukuoka, Fukuoka 812-8582, Japan				
TITLE	(E-mail:shinji@med.kyushu-u.ac.jp, Tel:81-92-642-5466,				
JOURNAL					
REFERENCE					
AUTHORS	Fax:81-92-642-5482)				
TITLE					
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AUTHORS	/translation="MEGAGCAGNKKKISSEKREKSRDAARSRSKSEVYELAHOL				
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KYTKDMEDIKILIASPSPHIHKETTSSTSPYDQTSRTSPNAGGIVETEOK  
SHPSRPNVLSVALSQRTPPEELPFLMLQNNQRKKMEHDSLFQAVGII"

BASE COUNT 1150 a 671 c 650 g 1080 t

ORIGIN

Query Match 93.6%; Score 23.4; DB 9; Length 3551;  
Best Local Similarity 96.0%; Pred. No. 0.38;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CCATGTGACCATTAAGAAATGAGAG 25  
|||||  
Db 440 CCATGTGACCATTAAGAAATGAGAG 464

RESULT 12  
HSU22431 3678 bp mRNA linear PRI 28-JUN-1995  
LOCUS  
DEFINITION Human hypoxia-inducible factor 1 alpha (HIF-1 alpha) mRNA, complete cds.  
ACCESSION U22431  
VERSION U22431.1 GI:881345  
KEYWORDS Homo sapiens.  
SOURCE Homo sapiens.  
ORGANISM Homo sapiens.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 3678)  
AUTHORS Wang, G.L., Jiang, B.-H., Rue, E.A. and Semenza, G.L.  
TITLE Hypoxia-inducible factor 1 is a basic-helix-loop-helix-PAS heterodimer regulated by cellular O2 tension  
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 92 (12), 5510-5514 (1995)  
MEDLINE 95296340  
PUBMED 7539918  
REFERENCE 2 (bases 1 to 3678)  
AUTHORS Wang, G.L., Jjiang, B.-H., Rue, E.A. and Semenza, G.L.  
TITLE Direct Submission  
JOURNAL Submitted (09-MAR-1995) Gregg L. Semenza, Center for Medical Genetics, The Johns Hopkins University School of Medicine, 600 N. Wolfe St., Baltimore, MD 21287-3914, USA

FEATURES  
source  
1..3678  
Location/Qualifiers  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/cell\_line="Hep3B"  
/cell\_type="hepatoblastoma"  
1..3678  
/gene="HIF-1 alpha"  
29..2509  
/gene="HIF-1 alpha"  
/standard\_name="hypoxia-inducible factor 1, alpha subunit"  
/note="basic helix-loop-helix transcription factor"  
/codon\_start=1  
/product="hypoxia-inducible factor 1 alpha"  
/protein\_id="AAC50152.1"  
/db\_xref="GI:881346"  
/translation="MEGAGAGANDKKKISSERRKESRDAAARRSRKSESEVEYELAHOL  
PLPHNVSHLDKASVMRLTISYLRKILGAGDDIDDDKAKMNCYFLKALDGFVAV  
LTDGDMYIISDNVNYKGLTGFELTGHVDFTHPCDHEMRMLTHRGVYKKGK  
ONTGSRFLRMKCTLTSGRTMNKISATWVLHCTGHIHYVDINSNOPOGKKPPMT  
CLVLCIEPIHPSPNIEIPLDSKTEFLSHSDMKESYCDERITELMGEPPELLGRSIV  
EYHALDSDLTHTHDMFTKQYTTQYMLAKRGYVWEVQATYITTKRSQOC  
IVCNVYVSGIIOHDLFSLQTEVCVLPVSSDMKTQLTVESDTSCLK  
EPDALTLAPAGDTIISLDGSDTEEDQOLEEPLYNVMPSPNELQMINIAM  
SPLEATPKPLRSSADPALNOEVALKEPNPSELSLFTMPDIODOTSPSGSTRO  
SSPEPNSPEYCEFDVSDMNEKLEVEKLFADTEPAKPFSTODDLLEMLAYI

PMDDPOLRSFOLSPSSASPESASPOSTVYFOQOIOEPTANATTTATTEL  
KYTKDMEDIKILIASPSPHIHKETTSSTSPYDQTSRTSPNAGGIVETEOK  
SHPSRPNVLSVALSQRTPPEELPFLMLQNNQRKKMEHDSLFQAVGIILOO  
PDDHATTSLSKRRYKCKSSQNMCKTIIILPSDLACLQSGMSDESLPOLTSY  
DEEVNAPIDGSSNLLQGEELLALQVN  
3678  
/gene="HIF-1 alpha"  
/note="42 A nucleotides"  
BASE COUNT 1197 a 695 c 675 g 1111 t

ORIGIN

Query Match 93.6%; Score 23.4; DB 9; Length 3678;  
Best Local Similarity 96.0%; Pred. No. 0.38;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CCATGTGACCATTAAGAAATGAGAG 25  
|||||  
Db 440 CCATGTGACCATTAAGAAATGAGAG 464

RESULT 13  
AF057308 3718 bp mRNA linear ROD 04-SEP-2001  
LOCUS  
DEFINITION Rattus norvegicus hypoxia-inducible factor-1 alpha (Hif1a) mRNA.  
ACCESSION AF057308  
VERSION AF057308.1 GI:4580532  
KEYWORDS Rattus norvegicus.  
SOURCE Rattus norvegicus.  
ORGANISM Rattus norvegicus.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
Rattus.  
REFERENCE 1 (bases 1 to 3718)  
AUTHORS Zou, A.P., Yang, Z.Z., Li, P.L. and Cowley, A.W. Jr.  
TITLE Oxygen-dependent expression of hypoxia-inducible factor-1alpha in renal medullary cells of rats  
JOURNAL Physiol. Genomics (Online) 6 (3), 159-168 (2001)  
MEDLINE 21417706  
PUBMED 11526200  
REFERENCE 2 (bases 1 to 3718)  
AUTHORS Zou, A.P., Su, N., Park, F., Li, P.L. and Cowley, A.W. Jr.  
TITLE Direct Submission  
JOURNAL Submitted (04-APR-1998) Departments of Physiology and Pharmacology, Medical College of Wisconsin, 8701 Watertown Plank Road, Milwaukee, WI 53226, USA

FEATURES  
source  
1..3718  
Location/Qualifiers  
/organism="Rattus norvegicus"  
/strain="Sprague Dawley"  
/db\_xref="taxon:10116"  
/sex="male"  
/tissue\_type="Kidney"  
/dev\_stage="adult"  
1..3718  
/gene="Hif1a"  
24..2495  
/gene="Hif1a"  
/note="transcription factor; RHIF-1"  
/codon\_start=1  
/product="hypoxia-inducible factor-1 alpha"  
/protein\_id="AAD24413.1"  
/db\_xref="GI:4580533"  
/translation="MEGAGAGENEKRNMSERRKESRDAAARRSRKSESEVEYELAHOL  
LPLPHNVSHLDKASVMRLTISYLRKILGAGDDIDDDKAKMNCYFLKALDGFVAV  
LTDGDMYIISDNVNYKGLTGFELTGHVDFTHPCDHEMRMLTHRGVYKKGK  
ONTGSRFLRMKCTLTSGRTMNKISATWVLHCTGHIHYVDINSNOPOGKKPPMT  
CLVLCIEPIHPSPNIEIPLDSKTEFLSHSDMKESYCDERITELMGEPPELLGRSIV  
EYHALDSDLTHTHDMFTKQYTTQYMLAKRGYVWEVQATYITTKRSQOC  
IVCNVYVSGIIOHDLFSLQTEVCVLPVSSDMKTQLTVESDTSCLK  
EPDALTLAPAGDTIISLDGSDTEEDQOLEEPLYNVMPSPNELQMINIAM  
SPLEATPKPLRSSADPALNOEVALKEPNPSELSLFTMPDIODOTSPSGSTRO  
SSPEPNSPEYCEFDVSDMNEKLEVEKLFADTEPAKPFSTODDLLEMLAYI



PMDDPOLRSEFOLSPLENSPSPSVTYTGFOQTOLOKPTTITVATATATDES  
 VTKNIEDIKILIASPSTOVPOEMTAKASATSGHSTRASPDAGKRVIEKTDKAH  
 PRSINLSTVLTQNTVPEEBELNPTLALQNAOKRKMEHDSLFQAAIGCTTLOQPD  
 RAPMSLSMKVKVGISSODGMEOKTIFLIPDLACRLGQMSDESGPOLTSYCE  
 VNAPIQSRNLQGEELLRALDQV"

BASE COUNT 1125 a 733 c 762 g 1098 t

Query Match 93.6%; Score 23.4; DB 10; Length 3718;  
 Best Local Similarity 96.0%; Pred. No. 0.38;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CCATGTGACCATAGGAATGAG 25  
 |||||  
 Db 438 CCATGTGACCATAGGAATGAG 462

## RESULT 14

AX306008 3746 bp DNA linear PAT 11-DEC-2001  
 LOCUS Sequence 759 from patent WO0188188.  
 DEFINITION AX306008  
 VERSION AX306008.1 GI:17645352  
 KEYWORDS  
 SOURCE house mouse.  
 ORGANISM Mus musculus.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1  
 AUTHORS Ishikawa, K., Asai, S., Takahashi, Y., Nagata, T. and Ishii, Y.  
 TITLE Method for examining ischemic conditions  
 JOURNAL Patent: WO 0188188-A 759 22-NOV-2001;  
 FEATURES School Juridical person Nihon University (JP)  
 Location/Qualifiers  
 Source 1..3746  
 /organism="Mus musculus"  
 /db\_xref="taxon:10090"

BASE COUNT 1124 a 740 c 740 g 1142 t  
 ORIGIN

Query Match 93.6%; Score 23.4; DB 6; Length 3746;  
 Best Local Similarity 96.0%; Pred. No. 0.38;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CCATGTGACCATAGGAATGAG 25  
 |||||  
 Db 501 CCATGTGACCATAGGAATGAG 525

RESULT 15  
 LOCUS MMHIF1ALP 3746 bp mRNA linear ROD 16-JUL-1996  
 DEFINITION M. musculus mRNA for hypoxia-inducible factor 1 alpha.  
 ACCESSION X95580  
 VERSION X95580.1 GI:1430864  
 KEYWORDS HIF-1alpha; hypoxia-inducible factor one alpha.  
 SOURCE Mus musculus.  
 ORGANISM Mus musculus.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 3746)  
 AUTHORS Wenger, R.H.  
 TITLE Direct Submission  
 JOURNAL Submitted (07-FEB-1996) R.H. Wenger, Institute of Physiology,  
 University of Zuerich-Irchel, Winterthurerstrasse 190, CH-8057  
 Zuerich, SWITZERLAND

REMARK Revised by author 16-JUL-96  
 2 (bases 1 to 3746)  
 REFERENCE Wenger, R.H., Rolfs, A., Marti, H.H., Guenet, J.L. and Gaasmann, M.  
 AUTHORS Nucleotide sequence, chromosomal assignment and mRNA expression of  
 TITLE mouse hypoxia-inducible factor-1 alpha  
 JOURNAL Biochem. Res. Commun. 223 (1), 54-59 (1996)  
 MEDLINE 96254028

PUBMED 8660378  
 COMMENT On Jul 17, 1996 this sequence version replaced gi:1419626.  
 RELATED sequence X95002 and U59496.  
 FEATURES Location/Qualifiers  
 source 1..3746

/organism="Mus musculus"  
 /db\_xref="taxon:10090"  
 /cell\_line="Hepatic7"  
 /cell\_type="hepatoma"  
 126..2558  
 /gene="HIF-1alpha"  
 126..2558  
 /gene="HIF-1alpha"  
 /codon\_start=1  
 /product="hypoxia-inducible factor one alpha"  
 /protein\_id="CAA64833.1"  
 /db\_xref="GI:4379202"  
 /db\_xref="MGI:106918"  
 /db\_xref="SWISS-PROT:O61221"  
 /translation="MSERRRKRSDAASRRKSEVRYELANQLPLPHVSSHLDK  
 ASVRLTISLYRRLKLDAGGSDSEDMKAQMDCTYLALDGFVAVLDDDMVYISD  
 NVKVMGLTQPELAGHSYDFTTHPCDHEEMRLNHRNGPVKRGKELNTQSFRLMK  
 CTTSGRTMNTKLSATWKVLCVHCTGHIHYVDITNSNPQCGYKRPMTCLVLCBPPIPH  
 SNTETPLDSKTEFLSRSLDMKFSYCDERTTELMGYEPBELGRSTIYEYVHMLDSDHIT  
 KTHHDMFTKGQVTTGQYRMLARKGGYVWEQATVITYNTRKNSQOCICVAVYVSGIT  
 OHDLIFSLQTESVYLKPYESSDMKMTOLFTEVESDITSCLPDKLKEPDALTLLAPA  
 GPTTISLDGSDDETEDQOLEDYLNDVMPSSNEKLNINLMSPLPSSSETPKPA  
 SSADPALNDEVALKLESSEBSLGISFTMPQIDDPASPSDSTQSSSPENSPSEYCP  
 DVSDDVMVNFLELEKLEAEDETEAKNPFSTQDDLDLEMLAPYIPMDDDPOLRSFPO  
 LSPLESNSPSPSMSTVTFQOTOLQKPTITATATATDESRTERRKDKDKEDIKILI  
 ASPSTQVPOQETITAKASAVSGTHSRTPRAGKRVIEOTDKAHPKINSATLNR  
 NTVPSEENPKTIASQNAOKRKMEHDSLFQAAIGCTTLOQPDCAPTMSLSMKRK  
 GFISSEONGTEBQKTIILIPDLACRLGQMSDVSGLPQLTSTDCENVAPIQGSKNLLQ  
 GEELLRALDQV"

BASE COUNT 1124 a 740 c 740 g 1142 t  
 ORIGIN

Query Match 93.6%; Score 23.4; DB 10; Length 3746;  
 Best Local Similarity 96.0%; Pred. No. 0.38;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CCATGTGACCATAGGAATGAG 25  
 |||||  
 Db 501 CCATGTGACCATAGGAATGAG 525

Search completed: March 17, 2003, 11:24:12  
 Job time : 236.108 secs





XX (ASPE-) ASPERA PHARM AB.  
 PA Poellinger L, Pereira T, Ruas J;  
 XX WPI; 2002-257466/30.  
 XX P-PSDB; AA077602.  
 DR New polypeptides comprising hypoxia-inducible factor-1 with alterations  
 PT of the transactivation domain, useful treating ischemic conditions,  
 PT e.g. brain infarction, heart infarction or circulatory disorder -  
 XX  
 PS Disclosure: Page 73-76; 80pp; English.  
 XX The invention relates to a polypeptide comprising hypoxia-inducible  
 CC factor-1 (HIF-1) with alterations of the transactivation domain (N-TAD or  
 CC C-TAD). Also included are nucleic acids encoding the altered proteins, a  
 CC vector comprising the nucleic acid, a host cell transformed with the  
 CC vector, methods for producing the protein or its functional fragment  
 CC or an isolated degradation box, a method of screening for an agent that  
 CC modulates N-TAD function and antagonists, agonists, modulators and  
 CC HIF-1 peptide fragments useful for modulating HIF-1 function or the  
 CC function of proteins that interact with it. The isolated polypeptides and  
 CC their fragments with altered residues are useful in methods for treating  
 CC diseases. The disease is an ischemic condition, e.g. brain infarction,  
 CC heart infarction or circulatory disorder. The disease may also be cancer,  
 CC hypertension, demyelinating disorders, diffuse proliferative  
 CC glomerulonephritis, toxoplasmosis caused retinochoroiditis, HIV (human  
 CC immunodeficiency virus) caused T4 angioedema, HIV-caused Kaposi's  
 CC sarcoma, hepatitis-caused inflammation, hepatitis-caused angiodysplasia,  
 CC chronic ulceration, proliferative retinopathy, retina haemangioblastomas,  
 CC neovascularisation, arterial hypervascularisation, sarcoidosis, bullous  
 CC skin disease, vasculitis with angioedema, dermatomyositis with  
 CC angioedema, polymyositis with angioedema, rheumatoid arthritis,  
 CC juvenile osteoarthritis, polyarthritis, aneurysm or atheroma. The  
 CC present sequence encodes HIF-1.  
 XX  
 SQ Sequence 2481 BP; 829 A; 512 C; 500 G; 640 T; 0 other;  
 Query Match 93.6%; Score 23.4; DB 24; Length 2481;  
 Best Local Similarity 96.0%; Pred. No. 0.19;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 CCATGTGACCATTTAGGAATGAGAG 25  
 |||||||||  
 DB 412 CCATGTGACCATTTAGGAATGAGAG 436  
 RESULT 2  
 ABL91695  
 ID ABL91695 standard; DNA: 2481 BP.  
 AC ABL91695;  
 XX 28-MAY-2002 (first entry)  
 DT  
 XX Human polynucleotide SEQ ID NO 38.  
 DE  
 XX Human; HIV; HCV; gene expression; oligonucleotide; tumour; pathogen;  
 KM Plasmidum; virus; viroid; cytokine; prion; antisense oligonucleotide;  
 KW Cytostatic; virucide; protozoacide; antibacterial; ds.  
 XX Homo sapiens.  
 OS  
 XX DE10100586-C1.  
 FN  
 XX 11-APR-2002.  
 PD  
 XX 09-JAN-2001; 2001DE-1000586.  
 PF  
 XX 09-JAN-2001; 2001DE-1000586.  
 PR  
 XX (RIBO-) RIBOPHARMA AG.

XX Kreutzer R, Limmer S, Rost S, Hadwiger P;  
 PI WPI; 2002-270454/32.  
 DR Inhibiting gene expression in cells, useful for e.g. treating tumours,  
 PT by introducing double-stranded complementary oligonucleotides having unpaired  
 PT terminal bases -  
 XX  
 PS Claim 13; Page 33; 104pp; German.  
 XX The invention relates to a method for inhibiting expression of a target  
 CC gene (ABL91658-ABL91797) in a cell by introducing at least one  
 CC oligonucleotide that has a double-stranded structure consisting of at  
 CC most 49 sequential nucleotide pairs, with at least part of one strand  
 CC complementary with the target gene and has at least one end a  
 CC single-stranded segment of 1-4 nt. The method provides  
 CC oligonucleotides for antisense inhibition of gene expression useful  
 CC e.g. for treating tumours but the oligonucleotides may also be  
 CC directed against genes present in pathogens (e.g. Plasmidum or  
 CC viruses/viroids, pathogenic on humans, animals or plants) or against  
 CC cytokine, Id, developmental or prion genes. The method provides more  
 CC effective inhibition of gene expression than use of known  
 CC oligonucleotides, probably because the unpaired overhang increases  
 CC stability and thus intracellular concentration.  
 XX  
 SQ Sequence 2481 BP; 829 A; 512 C; 500 G; 640 T; 0 other;  
 Query Match 93.6%; Score 23.4; DB 24; Length 2481;  
 Best Local Similarity 96.0%; Pred. No. 0.19;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 CCATGTGACCATTTAGGAATGAGAG 25  
 |||||||||  
 DB 412 CCATGTGACCATTTAGGAATGAGAG 436  
 RESULT 3  
 AAS14154  
 ID AAS14154 standard; CDNA: 2528 BP.  
 XX AAS14154;  
 AC 18-DEC-2001 (first entry)  
 DT  
 XX Human HIF-1 alpha DNA used in identification of hypoxia regulated genes.  
 DE  
 XX Differential expression; polymorphism; biological pathway; IRES; GTP; ss;  
 KM Internal ribosome entry site; green fluorescent protein; HIF-1 alpha;  
 KW hypoxia inducible factor 1 alpha; hypoxia regulated gene; macrophage;  
 KW human.  
 XX Homo sapiens.  
 OS  
 XX WO200162965-A2.  
 PN  
 XX 30-AUG-2001.  
 PD  
 XX 22-FEB-2001; 2001WO-GB00758.  
 PF  
 XX 22-FEB-2000; 2000GB-0004197.  
 PR  
 XX 28-JUL-2000; 2000GB-0016679.  
 XX (OXFO-) OXFORD BIOMEDICA UK LTD.  
 PA  
 XX Kingsman AJ;  
 PI  
 XX WPI; 2001-589807/66.  
 DR  
 XX Screening a genetic element involved in a cellular process, comprises  
 PT comparing gene expressions in a cell, and a second cell that has  
 PT altered levels of genes used in the process, and detecting an element  
 PT with varied expression -

XX PS Example 2: Page 96-97; 103pp; English.

CC The invention relates to differential expression screening to identify a  
CC genetic element involved in a cellular process. The method involves  
CC comparing gene expressions in two cells, where one cell has altered  
CC levels of a biological molecule, and identifying the genetic element  
CC whose expression differs, by comparing expression under different  
CC environmental conditions. The method is useful for identifying mutations  
CC and polymorphisms that affect the biological response to a particular  
CC cellular process. The method also allows the molecular dissection of  
CC biological pathways by altering a particular pathway under study. By  
CC introducing a heterologous nucleic acid into a cell population, the level  
CC of a particular molecule can be influenced, allowing a pathway to be  
CC dissected into its precise molecular components. The main use of the  
CC method is to compare gene expression in cells from a diseased patient and  
CC from a normal patient. This sequence represents a human hypoxia inducible  
CC factor 1 alpha (HIF-1 alpha) which is inserted into an Adapt Ires-GFP  
CC plasmid for use in identification of hypoxia-regulated genes in  
CC macrophages.

XX SQ Sequence 2528 BP; 839 A; 528 C; 513 G; 648 T; 0 other;

XX Query Match 93.6%; Score 23.4; DB 22; Length 2528;  
XX Best Local Similarity 96.0%; Pred. No. 0.19;  
XX Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATTTAGGAATGAGAG 25  
DB 439 CCATGTGACCATTTAGGAATGAGAG 463  
|||||

RESULT 4  
AAV63210  
ID AAV63210 standard; DNA; 3678 BP.  
XX AAV63210;  
XX  
XX AAV63210;  
XX  
XX 14-JAN-1999 (first entry)  
XX  
XX Nucleic acid sequence of human hypoxia inducible factor 1, alpha.  
XX  
XX Human transforming growth factor-beta 3; TGF-beta3; oxygen tension;  
XX trophoblast invasion regulation; inhibitor; HIF-1 alpha;  
XX TGF-beta family cytokine receptor; hypoxia inducible factor 1 alpha;  
XX preeclampsia; pregnancy; choriocarcinoma; ss.  
XX  
XX Homo sapiens.  
XX  
XX OS  
XX  
XX FH Key Location/Qualifiers  
XX FT CDS 29..2509  
XX FT /\*tag= a  
XX FT /product= HIF-1 alpha  
XX  
XX WO9840747-A1.  
XX  
XX 17-SEP-1998.  
XX  
XX PD  
XX  
XX PF 05-MAR-1998; 98WO-CA00180.  
XX  
XX PR 07-MAR-1997; 97US-0039919.  
XX  
XX  
XX (HOSP-) HOSPITAL FOR SICK CHILDREN.  
XX (MOUN) MOUNT SINAI HOSPITAL CORP.  
XX  
XX Canlgia I, Lye S, Post M;  
XX  
XX WPI; 1998-520837/44.  
XX P-PSDB; AAM80418.  
XX  
XX Regulation of trophoblast invasion - by, e.g. transforming growth  
XX factor-beta3 inhibitor, useful for detecting or treating  
XX preeclampsia in pregnant women

XX PS Disclosure; Fig 2; 59pp; English.

CC The present sequence encodes human hypoxia inducible factor 1 alpha  
CC (HIF-1 alpha). The specification describes a composition for regulating  
CC trophoblast invasion which comprises an inhibitor of transforming growth  
CC factor-beta 3 (TGF-beta3), TGF-beta family cytokine receptors, HIF-1  
CC alpha or oxygen tension. The composition is used in methods of  
CC diagnosing, monitoring, preventing or treating conditions requiring  
CC regulation of trophoblast invasion, especially preeclampsia in pregnant  
CC women or choriocarcinomas.

XX SQ Sequence 3678 BP; 1197 A; 695 C; 675 G; 1111 T; 0 other;

XX Query Match 93.6%; Score 23.4; DB 19; Length 3678;  
XX Best Local Similarity 96.0%; Pred. No. 0.2;  
XX Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATTTAGGAATGAGAG 25  
DB 440 CCATGTGACCATTTAGGAATGAGAG 464  
|||||

RESULT 5  
AAD38995  
ID AAD38995 standard; CDNA; 3678 BP.  
XX AAD38995;  
XX  
XX 23-SEP-2002 (first entry)  
XX  
XX Human HIF-1alpha CDNA.  
XX  
XX  
XX Human: haematologic malignancy; multidrug resistance; MDR; SUMO-1;  
XX hypoxia inducible factor-1; small ubiquitin-like modifier; HIF-1;  
XX lymphoid disorder; chronic lymphoproliferative disorder; lymphoma;  
XX myeloid disorder; lymphocytic leukaemia; thrombocythaemia; myeloma;  
XX angioendothelial myeloid metaplasia; myeloid leukaemia; gene therapy;  
XX polycythemia vera; hypoxia responsive element; HRE; gene; ss.  
XX  
XX Homo sapiens.  
XX  
XX OS  
XX  
XX FH Key Location/Qualifiers  
XX FT CDS 29..2509  
XX FT /\*tag= a  
XX FT /product= "Human HIF-1alpha protein"  
XX  
XX WO200234291-A2.  
XX  
XX 02-MAY-2002.  
XX  
XX PD 25-OCT-2001; 2001WO-US49856.  
XX  
XX PR 26-OCT-2000; 2000US-243542P.  
XX  
XX (BGHM) BRIGHAM & WOMENS HOSPITAL INC.  
XX  
XX Colgan SP;  
XX  
XX WPI; 2002-471427/50.  
XX P-PSDB; AAE24212.  
XX  
XX Treating a subject (at risk of) having a hematologic malignancy or  
XX multidrug resistance, e.g. lymphoma or myeloma, by administering  
XX hypoxia inducible factor 1 binding molecules or small  
XX ubiquitin-like-modifier-1 binding molecules -  
XX  
XX Disclosure; Page 61-63; 92pp; English.  
XX  
XX The invention relates to a method of treating a subject having or at  
XX risk of developing a haematologic malignancy or multidrug resistance  
XX (MDR). The method involves administering hypoxia inducible factor-1  
XX (HIF-1) binding molecules or small ubiquitin-like-modifier (SUMO)-1

CC binding molecules or HIF-1-SUMO-1 complex modulators. mdrl-hypoxia  
 CC responsive element (HRE) binding molecules or antisense nucleic  
 CC acid molecules and SUMO-1 binding molecules or antisense molecules  
 CC are useful for treating a subject having or at risk of developing  
 CC hematologic malignancy or MDR (e.g. a lymphoid disorder or a myeloid  
 CC disorder). The lymphoid disorders include lymphocytic leukemia or  
 CC chronic lymphoproliferative disorders e.g. lymphoma, myeloma or chronic  
 CC lymphoid leukemia. The myeloid disorders include chronic or acute  
 CC myeloid leukemia, e.g. angiotenic myeloid metaplasia, essential  
 CC thrombocythemia or polycythemia vera. The invention is used in gene  
 CC therapy. The present sequence is human HIF-1alpha cDNA.  
 CC  
 SQ Sequence 3678 BP; 1197 A; 695 C; 675 G; 1111 T; 0 other;  
 Query Match 93.6%; Score 23.4; DB 24; Length 3678;  
 Best Local Similarity 96.0%; Pred. No. 0.2;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 CCATGTGACCATAGGAATGAGAG 25  
 DB 440 CCATGTGACCATAGGAATGAGAG 464  
 RESULT 6  
 ABR84267  
 ID ABR84267 standard; cDNA; 3678 BP.  
 AC ABR84267;  
 XX  
 DT 14-AUG-2002 (first entry)  
 DE Human cDNA differentially expressed in granulocytic cells #838.  
 XX  
 KW Human; ss; granulocytic cell; DNA chip; bacterial infection;  
 KW viral infection; parasitic infection; protozoal infection;  
 KW fungal infection; sterile inflammatory disease; psoriasis;  
 KW rheumatoid arthritis; glomerulonephritis; asthma; thrombosis;  
 KW cardiac reperfusion injury; renal reperfusion injury; ARDS;  
 KW adult respiratory distress syndrome; inflammatory bowel disease;  
 KW Crohn's disease; ulcerative colitis; periodontal disease;  
 KW granulocyte activation; chronic inflammation; allergy.  
 KW  
 XX Homo sapiens.  
 OS  
 XX  
 PN WO200228999-A2.  
 XX  
 PD 11-APR-2002.  
 XX  
 PF 03-OCT-2001; 2001WO-US30821.  
 XX  
 PR 03-OCT-2000; 2000US-237189P.  
 XX  
 PA (GENE-) GENE LOGIC INC.  
 XX  
 PI Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;  
 XX  
 DR WPI: 2002-435328/46.  
 XX  
 XX  
 PT Detecting granulocyte activation by detecting differential expression  
 PT of genes associated with granulocyte activation, which serves as  
 PT diagnostic markers that is useful for monitoring disease states and  
 PT drug toxicity -  
 XX  
 PS Claim 1; SEQ ID No 838; 114pp; English.  
 XX  
 CC The invention relates to detecting (M1) granulocyte (GC) activation  
 CC (GCA), by detecting the level of expression of gene(s) (Gs) identified by  
 CC DNA chip analysis as given in the specification, and comparing  
 CC the expression level to an expression level in an unactivated  
 CC GC, where differential expression of Gs is indicative of GCA.  
 CC Also included are modulating (M2) GCA by contacting GC with an agent  
 CC that alters the expression of at least one gene in Gs; (2) screening (M3)  
 CC for an agent capable of modulating GCA or an inflammation (especially

CC chronic) in a tissue, an allergic response in a subject, exposure of a  
 CC subject to a pathogen or sterile inflammatory disease using the  
 CC gene expression profile; (3) detecting (M4) an inflammation (especially  
 CC chronic) in a tissue, an allergic response in a subject, exposure of a  
 CC subject to a pathogen or sterile inflammatory disease, by detecting the  
 CC level of expression in a sample of the tissue of gene(s) from Gs, where  
 CC the level of expression of the gene is indicative of inflammation;  
 CC (4) treating (M5) an inflammation (especially chronic) or in a tissue,  
 CC an allergic response in a subject, exposure of a subject to a pathogen  
 CC or sterile inflammatory disease, by contacting a tissue having  
 CC inflammation with an agent that modulates the expression of gene(s)  
 CC from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for  
 CC modulating GCA; M3 is useful for screening an agent capable of modulating  
 CC GCA preferably in an inflammation in a tissue; M4 is useful for  
 CC detecting an inflammation (especially chronic) in a tissue, an allergic  
 CC response in a subject, exposure of a subject to a pathogen or sterile  
 CC inflammatory disease (e.g. psoriasis, rheumatoid arthritis,  
 CC glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal  
 CC reperfusion injury, ARDS, adult respiratory distress syndrome,  
 CC inflammatory bowel disease, Crohn's disease, ulcerative colitis,  
 CC periodontal disease; also bacterial infection, viral infection,  
 CC parasitic infection, protozoal infection, fungal infection and M5 is  
 CC useful for treating one of the above conditions. The present  
 CC sequence represents a gene differentially expressed in granulocytes.  
 CC Note: The sequence data for this patent did not form part  
 CC of the printed specification, but was obtained in electronic  
 CC format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.  
 CC  
 SQ Sequence 3678 BP; 1197 A; 695 C; 675 G; 1111 T; 0 other;  
 Query Match 93.6%; Score 23.4; DB 24; Length 3678;  
 Best Local Similarity 96.0%; Pred. No. 0.2;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 CCATGTGACCATAGGAATGAGAG 25  
 DB 440 CCATGTGACCATAGGAATGAGAG 464  
 RESULT 7  
 AAT45937  
 ID AAT45937 standard; DNA; 3736 BP.  
 AC AAT45937;  
 XX  
 DT 19-MAR-1997 (first entry)  
 DE Human hypoxia inducible factor-1 alpha cDNA.  
 XX  
 DE Human hypoxia inducible factor-1 alpha cDNA.  
 XX  
 KW Hypoxia inducible factor-1 alpha; HIF-1; tissue damage;  
 KW atherosclerosis; cerebral artery disease; gene therapy; ss.  
 KW  
 XX Homo sapiens.  
 OS  
 XX  
 FH Key Location/Qualifiers  
 FH misc\_signal 23..32 /\*tag= a  
 FT /\*function= kozak sequence  
 FT CDS 29..2509 /\*tag= b  
 FT polyA\_signal 3674..3679 /\*tag= c  
 FT  
 XX  
 PN WO9639426-A1.  
 XX  
 PD 12-DEC-1996.  
 XX  
 PF 06-JUN-1996; 96WO-US10251.  
 XX  
 PR 06-JUN-1995; 95US-0480473.  
 XX  
 PA (UyGO ) UNIV JOHNS HOPKINS SCHOOL MED.

XX PI Semenaza GL;  
 XX DR WPI: 1997-043061/04.  
 XX P-PSDB; AAM06557.  
 PT DNA encoding human hypoxia-inducible factor 1 alpha - useful for  
 PT enhancing expression of structural gene and treatment of  
 XX hypoxia-related tissue damage  
 PS Claim 2; Page 49-53; 95pp; English.  
 CC A DNA sequence (AAT45937) codes for the alpha subunit (AAM06557) of  
 CC human hypoxia inducible factor-1 (HIF-1), a DNA binding protein  
 CC that binds to the enhancer region of e.g. erythropoietin and  
 CC vascular endothelial growth factor (VEGF) genes. The DNA sequence  
 CC is a composite of 3 clones obt. by screening an Hep3b library and  
 CC by database analysis. HIF-1 alpha in transformed host cells, as  
 CC prepare recombinant HIF-1 alpha in transfected host cells, as  
 CC probes, and in the gene therapy of HIF-1-mediated or hypoxia-related  
 CC disorders such as atherosclerotic coronary or cerebral artery  
 CC disease; antisense sequences inhibit HIF-1 expression e.g. to treat  
 CC tumour proliferation mediated by VEGF-induced angiogenesis.  
 SQ Sequence 3736 BP; 1255 A; 695 C; 675 G; 1111 T; 0 other;  
 Query Match 93.6%; Score 23.4; DB 18; Length 3736;  
 Best Local Similarity 96.0%; Pred. No. 0.2;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 OY 1 CCATGTGACCATTTAGGAATGAGAG 25  
 DB 440 CCATGTGACCATTTAGGAATGAGAG 464  
 RESULT 8  
 AA299537  
 ID AA299537 standard; DNA; 3736 BP.  
 AC AA299537;  
 XX 03-JUL-2000 (first entry)  
 DT  
 DE DNA encoding a wild type human hypoxia inducible factor-1 alpha.  
 XX  
 KW Human; hypoxia-inducible factor 1 alpha; HIF-1alpha; variant;  
 KW hypoxia inducible gene; hypoxia inducible factor; hypoxia;  
 KW ischemia related damage; angiogenesis; coronary artery disease;  
 KW ischemic tissue damage; ss.  
 OS Homo sapiens.  
 XX  
 XX Key Location/Qualifiers  
 FH 29.2509  
 FT CDS /tag= a  
 FT /product= "hypoxia inducible factor-1 alpha"  
 PN WO200010578-A1.  
 PD 02-MAR-2000.  
 XX  
 XX 25-AUG-1999; 99WO-US19416.  
 PF  
 XX 25-AUG-1998; 98US-0148547.  
 PR  
 XX (UYN0 ) UNIV JOHNS HOPKINS SCHOOL MEDICINE.  
 PA  
 PI Semenaza GL;  
 XX WPI: 2000-246493/21.  
 DR P-PSDB; AAY69407.  
 XX  
 PT Variant forms of hypoxia-inducible factor (HIF)-1 alpha, useful for

PT treating hypoxia or ischemia-related tissue damage -  
 XX  
 XX Disclosure; Page 80-89; 96pp; English.  
 PS  
 CC The present sequence encodes a wild type human hypoxia-inducible factor  
 CC (HIF)-1 alpha. The specification describes HIF-1alpha variants that are  
 CC stable under hypoxic and non-hypoxic conditions. The variants comprises  
 CC amino acid residues 1-391 and 521-826, 549-826, 576-826, 429-826,  
 CC 469-826, 494-826, 508-826, 512-826 or 517-826 of the wild type human  
 CC HIF-1alpha polypeptide, in which residues 551 and 552 are not serine  
 CC and threonine, respectively. The HIF-1alpha variant polynucleotide  
 CC sequences are useful for increasing expression of a hypoxia inducible  
 CC gene in a cell. They is also useful for providing constitutive  
 CC or preventing hypoxia or ischemia related damage. The variant  
 CC HIF-1alpha polypeptides are useful for providing prophylactic therapy  
 CC for inducing the level of angiogenesis in tissues of patients at risk  
 CC of coronary artery disease or ischemic tissue damage.  
 SQ Sequence 3736 BP; 1255 A; 695 C; 675 G; 1111 T; 0 other;  
 Query Match 93.6%; Score 23.4; DB 21; Length 3736;  
 Best Local Similarity 96.0%; Pred. No. 0.2;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 OY 1 CCATGTGACCATTTAGGAATGAGAG 25  
 DB 440 CCATGTGACCATTTAGGAATGAGAG 464  
 RESULT 9  
 AB199710  
 ID AB199710 standard; CDNA; 3746 BP.  
 XX  
 AC AB199710;  
 XX  
 DT 07-MAR-2002 (first entry)  
 DE  
 XX Mouse ischemic condition related CDNA sequence SEQ ID NO:759.  
 DE  
 XX Mouse; ischemia; compressive ischemia; occlusive ischemia;  
 KW vasospastic ischemia; ischemic condition; ischemic disease; ss.  
 XX  
 OS Mus musculus.  
 XX  
 PN WO200188188-A2.  
 XX  
 XX 22-NOV-2001.  
 PD  
 XX 18-MAY-2001; 2001WO-JF04192.  
 PF  
 XX 18-MAY-2000; 2000JP-0145977.  
 PR  
 XX (UYN1-) UNIV NIHON SCHOOL JURIDICAL PERSON.  
 PA  
 PI Ishikawa K, Asai S, Takehashi Y, Nagata T, Ishii Y;  
 XX  
 XX WPI: 2002-034733/04.  
 DR P-PSDB; ABB57270.  
 XX  
 XX Examining the ischemic condition (e.g. occlusive ischemia) by measuring  
 PT expression levels of particular genes defined in the specification or  
 PT by determining the expression profile of a gene group comprising these  
 PT genes -  
 XX  
 XX Claim 2; Page 1863-1869; 2690pp; English.  
 PS  
 XX The present invention describes a method for examining ischemic  
 CC conditions, comprising measuring the expression levels of particular  
 CC genes (I) in a test sample or determining the expression profile of a  
 CC gene group in the sample comprising genes selected from (I). The method  
 CC is useful for examining the ischemic condition (e.g. compressive  
 CC ischemia, occlusive ischemia or vasospastic ischemia) by measuring

CC expression levels of particular genes (AB199202 to AB199912, encoding  
 CC the protein sequences in AB57020 to AB57374) or by determining the  
 CC expression profile of a gene group comprising these genes. The  
 CC expression levels or expression profiles produced by these genes are  
 CC used as an indicator when screening for ischaemic condition-improving  
 CC drugs or therapeutics for ischaemic diseases. AB199913 and AB199914  
 CC represent PCR primers for a mouse ischaemic condition related sequence,  
 CC which are used in the exemplification of the present invention.

XX Sequence 3746 BP; 1124 A; 740 C; 740 G; 1142 T; 0 other;

Query Match 93.6%; Score 23.4; DB 24; Length 3746;  
 Best Local Similarity 96.0%; Pred. No. 0.2;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATGTAGGAATGAGAG 25  
 |||||  
 DB 501 CCATGTGACCATGTAGGAATGAGAG 525

RESULT 10  
 AAS61690  
 ID AAS61690 standard; cDNA: 3927 BP.  
 AC AAS61690;  
 XX  
 XX 29-JAN-2002 (first entry)  
 DT  
 XX Lung small cell carcinoma antigen, cDNA #231.  
 DE  
 XX Human; cytosolic; antitumour; lung small cell cancer antigen;  
 KW tumour; lung cancer; ss.  
 KM  
 XX Homo sapiens.  
 OS  
 XX MO200177168-A2.  
 PN  
 XX 18-OCT-2001.  
 PD  
 XX 11-APR-2001; 2001WO-US11859.  
 PF  
 XX 11-APR-2000; 2000US-196780P.  
 PR 21-JUN-2000; 2000US-213361P.  
 PR 01-SEP-2000; 2000US-229763P.  
 PR 05-SEP-2000; 2000US-230629P.  
 PR 14-SEP-2000; 2000US-232565P.  
 PR 19-DEC-2000; 2000US-257037P.  
 PR 08-JAN-2001; 2001US-260796P.  
 PA  
 XX (CORI-) CORIXA CORP.  
 PI  
 XX Lodes MJ, Wang T, Mohamath R, Indirias CY;  
 XX WPI: 2002-010896/01.  
 DR P-PSDB; AAU69409.  
 DR  
 XX Lung tumour polynucleotide and polypeptides useful in therapy and  
 PT diagnosis of cancer especially lung cancer -  
 XX  
 XX Claim 1; Page 201-202; 295pp; English.

XX The invention relates to novel isolated lung small cell cancer antigen  
 CC polynucleotides (I) and polypeptides (II) used in a method of detecting  
 CC cancer in a patient. The method is optionally performed by  
 CC utilising oligonucleotides (III), where the biological sample  
 CC from the patient is contacted with (III), detecting the amount of  
 CC polynucleotide hybridised to (III) in the sample and comparing the  
 CC amount of polynucleotide to a predetermined cut-off value and thereby  
 CC determining cancer in a patient. (I), (II) or antigen-presenting cells  
 CC expressing (II) is useful for stimulating and/or expanding T cells  
 CC specific for a tumour protein. The method comprises contacting T cells  
 CC with one of the components under conditions to permit the stimulation  
 CC and/or expansion of the cells. A composition comprising (I) is useful for

CC stimulating an immune response in a patient and for inhibiting the  
 CC development of a cancer especially lung cancer in a patient. An  
 CC isolated T cell population is useful for removing tumour cells from the  
 CC biological sample and for inhibiting the development of cancer in a  
 CC patient. AAS61460-AAS61874 represent novel human lung small cell  
 CC cancer antigen coding sequences of the invention.

XX Sequence 3927 BP; 1241 A; 782 C; 748 G; 1156 T; 0 other;

Query Match 93.6%; Score 23.4; DB 24; Length 3927;  
 Best Local Similarity 96.0%; Pred. No. 0.21;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATGTAGGAATGAGAG 25  
 |||||  
 DB 670 CCATGTGACCATGTAGGAATGAGAG 694

RESULT 11  
 AAX58980  
 ID AAX58980 standard; cDNA: 3933 BP.  
 AC AAX58980;  
 XX  
 XX 23-AUG-1999 (first entry)  
 DT  
 XX Human transcription regulator MOP1 cDNA.  
 DE  
 XX MOP1, member of the PAS superfamily; bHLH-PAS; human;  
 KW transcription regulator; hypoxia inducible factor 1 alpha; ss.  
 KM  
 XX Homo sapiens.  
 OS  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 265..2745  
 FT /\*tag= a  
 FT  
 XX MO9928464-A2.  
 PN  
 XX 10-JUN-1999.  
 PD  
 XX 27-NOV-1998; 98WO-US25314.  
 PF  
 XX 28-NOV-1997; 97US-0066863.  
 PR  
 XX (WISC ) WISCONSIN ALUMNI RES FOUND.  
 PA  
 XX Bradfield CA, Gu YZ, Hogenesch JB;  
 PI  
 XX WPI: 1999-371120/31.  
 DR P-PSDB; AAY06289.  
 DR  
 XX Developmental signal transduction associated proteins  
 PT  
 XX Example 1; Page 93-94; 106pp; English.

XX This is the nucleotide sequence of a cDNA encoding MOP1 (see  
 CC AAY06289), a member of the PAS superfamily, where PAS stands for  
 CC PER/ARNT/SIM domains. The cDNA was identified in an iterative  
 CC search of human ESTs designed to identify basic-helix-loop-helix-PAS  
 CC (bHLH-PAS) proteins that interact with either the Ah receptor (AHR)  
 CC or the Ah receptor nuclear translocator (ARNT). To obtain extended  
 CC open reading frames for each EST, an anchored-PCR strategy was used  
 CC to amplify additional flanking sequences from a commercial HepG2  
 CC library. MOP1 is also known as hypoxia inducible factor 1 alpha.  
 CC The factor is induced by low oxygen. It interacts with HSP90 and  
 CC with ARNT. The ARNT-dimerised factor regulates expression of  
 CC erythropoietin, among other genes. The invention also provides  
 CC novel MOPs 2-9 nucleic acids (see AAX58981-88) and proteins (see  
 CC AAY06289-97). These are useful in a variety of research,  
 CC diagnostic and therapeutic applications. Several of the MOPs are  
 CC alpha-class hypoxia-inducible factors. Others are involved in  
 CC circadian signal transduction.



XX Sequence 3933 BP; 1243 A; 784 C; 750 G; 1156 T; 0 other;  
SQ Query Match 93.6%; Score 23.4; DB 20; Length 3933;  
Best Local Similarity 96.0%; Pred. No. 0.21;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
Qy 1 CCATGTGACCATTTAGGAATGAGAG 25  
|||||  
Db 676 CCATGTGACCATTTAGGAATGAGAG 700

RESULT 12  
AAS85058  
ID AAS85058 standard; cDNA; 4162 BP.  
XX AAS85058;  
AC AAS85058;  
XX 13-FEB-2002 (first entry)  
DT  
XX DNA encoding novel human diagnostic protein #20862.  
DE  
XX Human; chromosome mapping; gene mapping; gene therapy; forensic;  
KM Food supplement; medical imaging; diagnostic; genetic disorder; ss.  
XX Homo sapiens.  
OS  
XX WO200175067-A2.  
PN  
XX 11-OCT-2001.  
PD  
XX 30-MAR-2001; 2001WO-US08631.  
PF  
XX 31-MAR-2000; 2000US-0540217.  
PR 23-AUG-2000; 2000US-0649167.  
XX (HYSE-) HYSEQ INC.  
PA  
PI Drmanac RT, Liu C, Tang YT;  
PI  
XX WPI; 2001-639362/73.  
DR P-PSDB; ABG20871.  
XX  
XX New isolated polynucleotide and encoded polypeptides, useful in  
PT diagnostics, forensics, gene mapping, identification of mutations  
PT responsible for genetic disorders or other traits and to assess  
PT biodiversity -  
XX  
XX Claim 1; SEQ ID NO 20862; 103pp; English.  
PS  
XX The invention relates to isolated polynucleotide (I) and  
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,  
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
CC and gene mapping, and in recombinant production of (II). The  
CC polynucleotides are also used in diagnostics as expressed sequence tags  
CC for identifying expressed genes. (I) is useful in gene therapy techniques  
CC to restore normal activity of (II) or to treat disease states involving  
CC (II). (II) is useful for generating antibodies against it, detecting or  
CC quantifying a polypeptide in tissue, as molecular weight markers and as  
CC a food supplement. (II) and its binding partners are useful in medical  
CC imaging of sites expressing (II). (I) and (II) are useful for treating  
CC disorders involving aberrant protein expression or biological activity.  
CC The polypeptide and polynucleotide sequences have applications in  
CC diagnostics, forensics, gene mapping, identification of mutations  
CC responsible for genetic disorders or other traits to assess biodiversity  
CC and to produce other types of data and products dependent on DNA and  
CC amino acid sequences. AAS64197-AAS94564 represent novel human  
CC diagnostic coding sequences of the invention.  
CC Note: The sequence data for this patent did not appear in the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at [http://wipo.int/publ/published\\_pcc\\_sequences](http://wipo.int/publ/published_pcc_sequences).  
XX  
XX Sequence 4162 BP; 1286 A; 843 C; 813 G; 1220 T; 0 other;

Query Match 93.6%; Score 23.4; DB 23; Length 4162;  
Best Local Similarity 96.0%; Pred. No. 0.21;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
Qy 1 CCATGTGACCATTTAGGAATGAGAG 25  
|||||  
Db 804 CCATGTGACCATTTAGGAATGAGAG 828

RESULT 13  
AAS14156  
ID AAS14156 standard; DNA; 10355 BP.  
XX AAS14156;  
AC AAS14156;  
XX 18-DEC-2001 (first entry)  
DT  
XX PSMARF CMV-HIF DNA from a vector expressing HIF-1 alpha.  
DE  
XX Differential expression; polymorphism; biological pathway; IRS; GFP; ds;  
KM Internal ribosome entry site; green fluorescent protein; HIF-1 alpha;  
KM hypoxia inducible factor 1 alpha; hypoxia regulated gene; macrophage;  
KM human; CMV; cytomegalovirus.  
XX Synthetic.  
OS  
XX WO200162965-A2.  
PN  
XX 30-AUG-2001.  
PD  
XX 22-FEB-2001; 2001WO-GB00758.  
PF  
XX 22-FEB-2000; 2000GB-0004197.  
PR 28-JUL-2000; 2000GB-0018679.  
XX (OXFO-) OXFORD BIOMEDICA UK LTD.  
PA  
PI Kingsman AJ;  
PI  
XX WPI; 2001-589807/66.  
DR  
XX  
XX Screening a genetic element involved in a cellular process, comprises  
PT comparing gene expressions in a cell, and a second cell that has  
PT altered levels of genes used in the process, and detecting an element  
PT with varied expression -  
XX  
XX Example 5; Page 97-101; 103pp; English.  
PS  
XX The invention relates to differential expression screening to identify a  
CC genetic element involved in a cellular process. The method involves  
CC comparing gene expressions in two cells, where one cell has altered  
CC levels of a biological molecule, and identifying the genetic element  
CC whose expression differs, by comparing expression under different  
CC environmental conditions. The method is useful for identifying mutations  
CC and polymorphisms that affect the biological response to a particular  
CC cellular process. The method also allows the molecular dissection of  
CC biological pathways by altering a particular pathway under study. By  
CC introducing a heterologous nucleic acid into a cell population, the level  
CC of a particular molecule can be influenced, allowing a pathway to be  
CC dissected into its precise molecular components. The main use of the  
CC method is to compare gene expression in cells from a diseased patient and  
CC from a normal patient. This sequence represents PSMARF CMV-HIF from a  
CC vector expressing hypoxia inducible factor 1 alpha (HIF-1 alpha) which is  
CC used in methods of the invention.  
XX  
XX Sequence 10355 BP; 3060 A; 2212 C; 2373 G; 2710 T; 0 other;  
SQ  
Query Match 93.6%; Score 23.4; DB 22; Length 10355;  
Best Local Similarity 96.0%; Pred. No. 0.24;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
Qy 1 CCATGTGACCATTTAGGAATGAGAG 25

DB 3305 CCATGTGACATGACGAAATGAGAG 3329  
|||||

## RESULT 14

AAK7781/C

ID AAK7781 standard; DNA: 27884 BP.

AC AAK7781;

DT 07-NOV-2001 (first entry)

XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:32593.

DE Human immune/haematopoietic; immune/haematopoietic antigen; cancer;

KW cytostatic; gene therapy; vaccine; metastasis; ds.

XX Homo sapiens.

XX WO200157182-A2.

XX 09-AUG-2001.

XX 17-JAN-2001; 2001WO-US01354.

XX 31-JAN-2000; 2000US-0179065.

PR 04-FEB-2000; 2000US-0180628.

PR 24-FEB-2000; 2000US-0184564.

PR 02-MAR-2000; 2000US-0186350.

PR 16-MAR-2000; 2000US-0189874.

PR 17-MAR-2000; 2000US-0190076.

PR 18-APR-2000; 2000US-0198123.

PR 19-MAY-2000; 2000US-0205515.

PR 07-JUN-2000; 2000US-0209467.

PR 28-JUN-2000; 2000US-0214886.

PR 30-JUN-2000; 2000US-0215133.

PR 07-JUL-2000; 2000US-0216647.

PR 07-JUL-2000; 2000US-0216880.

PR 11-JUL-2000; 2000US-0217487.

PR 11-JUL-2000; 2000US-0217496.

PR 14-JUL-2000; 2000US-0218290.

PR 26-JUL-2000; 2000US-0220963.

PR 26-JUL-2000; 2000US-0220964.

PR 14-AUG-2000; 2000US-0224518.

PR 14-AUG-2000; 2000US-0224519.

PR 14-AUG-2000; 2000US-0225213.

PR 14-AUG-2000; 2000US-0225214.

PR 14-AUG-2000; 2000US-0225266.

PR 14-AUG-2000; 2000US-0225267.

PR 14-AUG-2000; 2000US-0225268.

PR 14-AUG-2000; 2000US-0225270.

PR 14-AUG-2000; 2000US-0225447.

PR 14-AUG-2000; 2000US-0225457.

PR 08-SEP-2000; 2000US-0231414.  
PR 08-SEP-2000; 2000US-0232080.  
PR 08-SEP-2000; 2000US-0232081.  
PR 12-SEP-2000; 2000US-0231968.  
PR 14-SEP-2000; 2000US-0232397.  
PR 14-SEP-2000; 2000US-0232398.  
PR 14-SEP-2000; 2000US-0232399.  
PR 14-SEP-2000; 2000US-0232400.  
PR 14-SEP-2000; 2000US-0232401.  
PR 14-SEP-2000; 2000US-0233063.  
PR 14-SEP-2000; 2000US-0233064.  
PR 14-SEP-2000; 2000US-0233065.  
PR 21-SEP-2000; 2000US-0234223.  
PR 21-SEP-2000; 2000US-0234274.  
PR 25-SEP-2000; 2000US-0234597.  
PR 25-SEP-2000; 2000US-0234598.  
PR 25-SEP-2000; 2000US-0234599.  
PR 26-SEP-2000; 2000US-0235484.  
PR 27-SEP-2000; 2000US-0235834.  
PR 27-SEP-2000; 2000US-0235836.  
PR 29-SEP-2000; 2000US-0236327.  
PR 29-SEP-2000; 2000US-0236327.  
PR 29-SEP-2000; 2000US-0236367.  
PR 29-SEP-2000; 2000US-0236368.  
PR 29-SEP-2000; 2000US-0236369.  
PR 29-SEP-2000; 2000US-0236370.  
PR 02-OCT-2000; 2000US-0236802.  
PR 02-OCT-2000; 2000US-0237037.  
PR 02-OCT-2000; 2000US-0237038.  
PR 02-OCT-2000; 2000US-0237039.  
PR 02-OCT-2000; 2000US-0237040.  
PR 13-OCT-2000; 2000US-0239335.  
PR 13-OCT-2000; 2000US-0239337.  
PR 20-OCT-2000; 2000US-0240960.  
PR 20-OCT-2000; 2000US-0241221.  
PR 20-OCT-2000; 2000US-0241785.  
PR 20-OCT-2000; 2000US-0241786.  
PR 20-OCT-2000; 2000US-0241787.  
PR 20-OCT-2000; 2000US-0241808.  
PR 20-OCT-2000; 2000US-0241809.  
PR 20-OCT-2000; 2000US-0241826.  
PR 01-NOV-2000; 2000US-0244617.  
PR 08-NOV-2000; 2000US-0244674.  
PR 08-NOV-2000; 2000US-0244675.  
PR 08-NOV-2000; 2000US-0244676.  
PR 08-NOV-2000; 2000US-0244677.  
PR 08-NOV-2000; 2000US-0244678.  
PR 08-NOV-2000; 2000US-0244679.  
PR 08-NOV-2000; 2000US-0244680.  
PR 08-NOV-2000; 2000US-0244681.  
PR 08-NOV-2000; 2000US-0244682.  
PR 08-NOV-2000; 2000US-0244683.  
PR 08-NOV-2000; 2000US-0244684.  
PR 08-NOV-2000; 2000US-0244685.  
PR 08-NOV-2000; 2000US-0244686.  
PR 08-NOV-2000; 2000US-0244687.  
PR 08-NOV-2000; 2000US-0244688.  
PR 08-NOV-2000; 2000US-0244689.  
PR 08-NOV-2000; 2000US-0244690.  
PR 08-NOV-2000; 2000US-0244691.  
PR 08-NOV-2000; 2000US-0244692.  
PR 08-NOV-2000; 2000US-0244693.  
PR 08-NOV-2000; 2000US-0244694.  
PR 08-NOV-2000; 2000US-0244695.  
PR 08-NOV-2000; 2000US-0244696.  
PR 08-NOV-2000; 2000US-0244697.  
PR 08-NOV-2000; 2000US-0244698.  
PR 08-NOV-2000; 2000US-0244699.  
PR 08-NOV-2000; 2000US-0244700.  
PR 08-NOV-2000; 2000US-0244701.  
PR 08-NOV-2000; 2000US-0244702.  
PR 08-NOV-2000; 2000US-0244703.  
PR 08-NOV-2000; 2000US-0244704.  
PR 08-NOV-2000; 2000US-0244705.  
PR 08-NOV-2000; 2000US-0244706.  
PR 08-NOV-2000; 2000US-0244707.  
PR 08-NOV-2000; 2000US-0244708.  
PR 08-NOV-2000; 2000US-0244709.  
PR 08-NOV-2000; 2000US-0244710.  
PR 08-NOV-2000; 2000US-0244711.  
PR 08-NOV-2000; 2000US-0244712.  
PR 08-NOV-2000; 2000US-0244713.  
PR 08-NOV-2000; 2000US-0244714.  
PR 08-NOV-2000; 2000US-0244715.  
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PR 08-NOV-2000; 2000US-0244718.  
PR 08-NOV-2000; 2000US-0244719.  
PR 08-NOV-2000; 2000US-0244720.  
PR 08-NOV-2000; 2000US-0244721.  
PR 08-NOV-2000; 2000US-0244722.  
PR 08-NOV-2000; 2000US-0244723.  
PR 08-NOV-2000; 2000US-0244724.  
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PR 08-NOV-2000; 2000US-0244726.  
PR 08-NOV-2000; 2000US-0244727.  
PR 08-NOV-2000; 2000US-0244728.  
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PR 08-NOV-2000; 2000US-0244730.  
PR 08-NOV-2000; 2000US-0244731.  
PR 08-NOV-2000; 2000US-0244732.  
PR 08-NOV-2000; 2000US-0244733.  
PR 08-NOV-2000; 2000US-0244734.  
PR 08-NOV-2000; 2000US-0244735.  
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PR 08-NOV-2000; 2000US-0244738.  
PR 08-NOV-2000; 2000US-0244739.  
PR 08-NOV-2000; 2000US-0244740.  
PR 08-NOV-2000; 2000US-0244741.  
PR 08-NOV-2000; 2000US-0244742.  
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PR 08-NOV-2000; 2000US-0244746.  
PR 08-NOV-2000; 2000US-0244747.  
PR 08-NOV-2000; 2000US-0244748.  
PR 08-NOV-2000; 2000US-0244749.  
PR 08-NOV-2000; 2000US-0244750.  
PR 08-NOV-2000; 2000US-0244751.  
PR 08-NOV-2000; 2000US-0244752.  
PR 08-NOV-2000; 2000US-0244753.  
PR 08-NOV-2000; 2000US-0244754.  
PR 08-NOV-2000; 2000US-0244755.  
PR 08-NOV-2000; 2000US-0244756.  
PR 08-NOV-2000; 2000US-0244757.  
PR 08-NOV-2000; 2000US-0244758.  
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PR 08-NOV-2000; 2000US-0244760.  
PR 08-NOV-2000; 2000US-0244761.  
PR 08-NOV-2000; 2000US-0244762.  
PR 08-NOV-2000; 2000US-0244763.  
PR 08-NOV-2000; 2000US-0244764.  
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PR 08-NOV-2000; 2000US-0244766.  
PR 08-NOV-2000; 2000US-0244767.  
PR 08-NOV-2000; 2000US-0244768.  
PR 08-NOV-2000; 2000US-0244769.  
PR 08-NOV-2000; 2000US-0244770.  
PR 08-NOV-2000; 2000US-0244771.  
PR 08-NOV-2000; 2000US-0244772.  
PR 08-NOV-2000; 2000US-0244773.  
PR 08-NOV-2000; 2000US-0244774.  
PR 08-NOV-2000; 2000US-0244775.  
PR 08-NOV-2000; 2000US-0244776.  
PR 08-NOV-2000; 2000US-0244777.  
PR 08-NOV-2000; 2000US-0244778.  
PR 08-NOV-2000; 2000US-0244779.  
PR 08-NOV-2000; 2000US-0244780.  
PR 08-NOV-2000; 2000US-0244781.  
PR 08-NOV-2000; 2000US-0244782.  
PR 08-NOV-2000; 2000US-0244783.  
PR 08-NOV-2000; 2000US-0244784.  
PR 08-NOV-2000; 2000US-0244785.  
PR 08-NOV-2000; 2000US-0244786.  
PR 08-NOV-2000; 2000US-0244787.  
PR 08-NOV-2000; 2000US-0244788.  
PR 08-NOV-2000; 2000US-0244789.  
PR 08-NOV-2000; 2000US-0244790.  
PR 08-NOV-2000; 2000US-0244791.  
PR 08-NOV-2000; 2000US-0244792.  
PR 08-NOV-2000; 2000US-0244793.  
PR 08-NOV-2000; 2000US-0244794.  
PR 08-NOV-2000; 2000US-0244795.  
PR 08-NOV-2000; 2000US-0244796.  
PR 08-NOV-2000; 2000US-0244797.  
PR 08-NOV-2000; 2000US-0244798.  
PR 08-NOV-2000; 2000US-0244799.  
PR 08-NOV-2000; 2000US-0244800.

PA (HUMA-) HUMAN GENOME SCI INC.  
XX  
PI Rosen CA, Barash SC, Ruben SM,  
XX  
DR WPI; 2001-483426/52.  
XX

Disclosure; SEQ ID NO 32593; 3071pp + Sequence Listing; English.

Sequence 27884 BP; 9296 A; 5084 C; 5260 G; 8244 T; 0 other;

Query Match	93.68;	Score 23.4;	DB 22;	Length 27884;
Best Local Similarity	96.04;	Pred. No. 0.27;		
Matches 24;	Conservative 0;	Mismatches 1;	Indels 0;	Gaps 0;

Y 1 CCATGTGACCATTAGGAAATGAGAG 25  
|||||  
b 26482 CCATGTGACCATGAGGAATGAGAG 26458

Accession	Gene	Product	Length (bp)	Accession	Gene	Product	Length (bp)
AAH93977	CDNA	367	BP.	AAH93977	CDNA	367	BP.
05-OCT-2001	(first entry)			05-OCT-2001	(first entry)		
Human foetal CDNA, SEQ ID NO: 506.				Human foetal CDNA, SEQ ID NO: 506.			
Human: foetal protein; cytosolic; immunosuppressive; immunostimulant; neuroprotective; thrombolytic; osteopathic; antiinflammatory; gene therapy; antisense therapy; cancer; immune disorder; growth disorder; osteoporosis; thrombolytic disorder; nervous system disorder; inflammation; expressed sequence tag; EST; ss.				Human: foetal protein; cytosolic; immunosuppressive; immunostimulant; neuroprotective; thrombolytic; osteopathic; antiinflammatory; gene therapy; antisense therapy; cancer; immune disorder; growth disorder; osteoporosis; thrombolytic disorder; nervous system disorder; inflammation; expressed sequence tag; EST; ss.			

PN WO200155339-A2.  
XX

25-JAN-2001; 2001WO-US02723.

PR 25-JAN-2000; 2000US-0491404.  
PR 15-SEP-2000; 2000US-0663870

PR 06-NOV-2000; 2000US-0707351.  
XX

PA (HYSE-) HYSEQ INC.  
XX

PI Yeung G, Ford JE, Boyle BJ, Arterburn MC, Dimanac RA, Tang YT;  
PI Liu C, Asundi V, Zhou P, Werhman T;  
XX

DR WPI; 2001-465571/50.  
DR P-PSDB; AAM06302.  
XX

PT Novel fetal proteins useful for the treatment and diagnosis of diseases associated with dysfunction of the protein e.g. cancers, immune disorders, growth disorders, thrombolytic disorders, nervous system disorders and inflammation -

Claim 1; Page 362-363; 715pp; English.

The invention relates to novel foetal polypeptides encoded by polynucleotides comprising one of 477 sequences fully defined in the specification. The foetal polynucleotides and polypeptides are useful in the treatment and diagnosis of diseases such as cancers, immune disorders, growth disorders (e.g. osteoporosis), thrombolytic disorders, nervous system disorders and inflammation. The present sequence was assembled using an expressed sequence tag (EST) found to be expressed in human foetal tissue cDNA libraries as the seed.

Query Match	72.8%	Score 18.2;	DB 22;	Length 367;
Best Local Similarity	87.0%;	Pred. No. 44;		
Matches 20;	Conservative 0;	Mismatches 3;	Indels 0;	Gaps 0

3 ATGTGACCATTAGGAATGAGAG 25  
 219 ATGTGACCATTAGGAATGAGAG 197

Search completed: March 17, 2003, 10:50:36  
Job time : 151.446 secs



GenCore version 5.1.4\_p5\_4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 10:23:12 ; Search time 883.737 Seconds  
(without alignments)  
458.154 Million cell updates/sec

Title: US-09-836-439-2

Perfect score: 25  
Sequence: 1 ccagtgaccattaggaatgagag 25

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 809774376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database :

EST:\*  
1: em\_estda:\*  
2: em\_esthum:\*  
3: em\_estlin:\*  
4: em\_estlun:\*  
5: em\_estlov:\*  
6: em\_estlpl:\*  
7: em\_estro:\*  
8: em\_hlc:\*  
9: em\_hlc1:\*  
10: em\_hlc2:\*  
11: em\_hlc3:\*  
12: em\_hlc4:\*  
13: em\_hlc5:\*  
14: em\_hlc6:\*  
15: em\_hlc7:\*  
16: em\_hlc8:\*  
17: em\_hlc9:\*  
18: em\_hlc10:\*  
19: em\_hlc11:\*  
20: em\_hlc12:\*  
21: em\_hlc13:\*  
22: em\_hlc14:\*  
23: em\_hlc15:\*  
24: em\_hlc16:\*  
25: em\_hlc17:\*  
26: em\_hlc18:\*  
27: em\_hlc19:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match Length	ID	Description
1	23.4	93.6	337	BQ308862 MR0-BT550
2	23.4	93.6	340	AM504525 UI-HF-BNO
3	23.4	93.6	380	AM120791 UI-M-BH2
4	23.4	93.6	386	AM124493 UI-M-BH2
5	23.4	93.6	395	BE980521 UI-M-BG2
6	23.4	93.6	501	BF406698 UI-R-BJ2

Result No.	Score	Query Match Length	ID	Description
7	23.4	93.6	591	AI481068 v192d03.x
8	23.4	93.6	684	BM834484 K-EST0109
9	23.4	93.6	693	BB199902 BB199902
10	23.4	93.6	711	BE731662 601567048
11	23.4	93.6	734	BF789080 602104940
12	23.4	93.6	739	BI413382 602296733
13	23.4	93.6	769	BG772697 602720844
14	23.4	93.6	791	BG110077 602279683
15	23.4	93.6	878	AI132294 AI132294
16	23.4	93.6	878	AI132294 AI132294
17	23.4	93.6	889	BQ213144 AGENCOURT
18	23.4	93.6	893	BE877244 AGENCOURT
19	23.4	93.6	941	BQ881947 AGENCOURT
20	23.4	93.6	1024	BM807153 AGENCOURT
21	23.4	93.6	1657	AK017853 muscu
22	22.4	89.6	714	AI131767 AI131767
23	21.8	87.2	310	BB204614 BB204614
24	21.8	87.2	903	AL698195 DKFZP6860
25	20.4	81.6	533	A2786928 2M0032N05
26	19.8	79.2	372	W86995 zh61e02.81
27	19.8	79.2	478	AA700482 z174f07.s
28	19.8	79.2	1435	BM906863 AGENCOURT
29	19.2	76.8	105	BQ308421 MR0-BT450
30	19.2	76.8	379	T31654 EST36504 Hu
31	19.2	76.8	415	AI0108979 CIT-HSP-2
32	19.2	76.8	611	B90780 CIT-HSP-217
33	18.8	75.2	381	A2263188 RPT-23-1
34	18.8	75.2	632	BQ397883 NISC mo02
35	18.8	75.2	632	BB652302 BB652302
36	18.8	75.2	800	BF617665 HSMC001
37	18.6	74.4	260	A2778470 2M0018K17
38	18.6	74.4	335	BJ198294 BJ198294
39	18.6	74.4	366	A2433190 1M0218L22
40	18.6	74.4	377	BJ195711 BJ195711
41	18.6	74.4	425	A2931901 474.dhz91
42	18.6	74.4	468	AL184121 Telradon
43	18.6	74.4	484	AA203974 mu28905.r
44	18.6	74.4	519	BM482542 535462 MA
45	18.6	74.4	538	BH026082 RPT-24-3

## ALIGNMENTS

RESULT 1  
LOCUS BQ308862 337 bp mRNA linear EST 16-MAY-2002  
DEFINITION MR0-BT5505-040701-003-c02 BT5505 Homo sapiens CDNA, mRNA sequence.  
ACCESSION BQ308862  
VERSION BQ308862.1 GI:20851208  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE 1 (bases 1 to 337)  
Nagai M.A., de Silva M.Jr., Zago M.A., Bordin S., Costa F.F., Goldman G.H., Carvalho A.F., Matsukuma A., Bala G.S., Simpson D.H., Brunstein A., de Oliveira P.S., Bucher P., Jongeneel C.V., O'Hare M.J., Soares F., Brentani R.R., Reis L.F., de Souza S.J. and Simpson A.J.  
Shotgun sequencing of the human transcriptome with ORF expressed sequence tags  
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001

Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL:  
(http://www.ludwig.org.br/scripts/gethtml2.pl?l1=MR0&l2=MR0-BT5505-040701-003-c02&l3=2001-07-04&l4=1)

Seq primer: puc 18 forward  
High quality sequence start: 15  
High quality sequence stop: 331.  
Location/Qualifiers

## FEATURES

source

1. 337

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone\_lib="BT5505"

/dev\_stage="Adult"

/note="Organ: breast; Vector: puc18; Site:1; Sma1; Site:2; Sma1; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196 716 - Ludwig Institute for Cancer Research) profiles into the puc 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

BASE COUNT 105 a 49 c 81 g 102 t  
ORIGIN

Query Match 93.6%; Score 23.4; DB 14; Length 337;  
Best Local Similarity 96.0%; Pred. No. 2.7;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCATGTGACCATTTAGGAATGAGAG 25  
|||||

Db 259 CCATGTGACCATTTAGGAATGAGAG 283  
|||||

RESULT 2 348 bp mRNA linear EST 02-MAR-2000  
AM504525  
LOCUS  
DEFINITION  
IMAGE:3079736 5', mRNA sequence.  
AM504525  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
Homo sapiens  
human.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 348)  
NIH-MGC http://mgi.nci.nih.gov/  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: cgapds@mail.nih.gov  
Eco RI site shown at the beginning of the sequence.  
Tissue Procurement: Louis M. Staudt, M.D., Ph.D.  
cDNA Library Preparation: M.B. Soares Lab  
cDNA Library Arrayed by: M.B. Soares Lab  
DNA Sequencing by: M.B. Soares Lab  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:  
www.bio.llnl.gov/dbp/image/image.html  
Seq primer: M13 Forward.

## FEATURES

source

Location/Qualifiers

1. 348

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone\_image="3079736"

/clone\_lib="NIH MGC 50"

/tissue\_type="lymph"

/cell\_type="germinal center B cells"

/cell\_line="MGC85"

/lab\_host="DH10B (LT1)"

/note="Vector: pT73-Pac; Site:1; NotI; Site:2; Eco RI; constructed from size fractionated cytoplasmic mRNA (3.5-4.4kb). Directionally cloned. Cells provided by

## BASE COUNT

107 a 54 c 81 g 106 t  
ORIGIN

Query Match 93.6%; Score 23.4; DB 10; Length 348;  
Best Local Similarity 96.0%; Pred. No. 2.7;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCATGTGACCATTTAGGAATGAGAG 25  
|||||

Db 255 CCATGTGACCATTTAGGAATGAGAG 279  
|||||

RESULT 3 380 bp mRNA linear EST 22-OCT-1999  
AM120791/c  
LOCUS  
DEFINITION  
IMAGE:3079736 5', mRNA sequence.  
AM120791  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
Mus musculus  
house mouse.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 380)  
Normalizaton and subtraction: two approaches to facilitate gene discovery  
Genome Res. 6 (9), 791-806 (1996)  
JOURNAL  
MEDLINE  
COMMENT  
Contact: Chin, H  
National Institute of Mental Health  
6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD  
20892-9643, USA  
Tel: 301 443 1706  
Fax: 301 443 9890  
Email: mestr@mail.nih.gov  
The sequence contained an oligo-dT track that was present in the oligonucleotide that was used to prime the synthesis of first strand cDNA and therefore this may represent a bonafide poly A tail. The sequence tag present in the cDNA between the NotI site and the oligo-dT track served to identify it as a clone from the normalized basal ganglia library cDNA library Preparation: M.B. Soares Lab clone distribution: NIH BMAP cDNA clones will be made available by the means that is soon to be determined. When NIH determines the means for distribution of the BMAP cDNA clones, this record will be updated accordingly when that means is determined.  
Seq primer: M13 Forward  
POLYA=Yes.

## FEATURES

source

Location/Qualifiers

1. 380

/organism="Mus musculus"

/strain="C57BL/6J"

/db\_xref="taxon:10090"

/clone="UI-M-BH2.3-any-h-05-0-UI"

/clone\_lib="NIH BMAP M.S3.3"

/dev\_stage="27-32 days"

/lab\_host="DH10B (Life Technologies)"

/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site:1; Not I; Site:2; Eco RI; The NIH\_BMAP\_M.S3.3 library is a subtracted library of a series, ultimately derived from a mixture of individually tagged normalized libraries from ten regions of the mouse brain (cerebellum, brain stems, olfactory bulbs, hypothalamus, cortex, amygdala, basal ganglia, pineal gland, striatum, hippocampus) after a series of subtractions to reduce the representation of cDNAs from which ESTs had already been generated. The following serially subtracted libraries were generated in this process: NIH\_BMAP\_M.S3.3, NIH\_BMAP\_M.S2, NIH\_BMAP\_M.S1. The subtracted library (NIH\_BMAP\_M.S3.3) was constructed

as follows: PCR amplified cDNA inserts from NIH\_BMAP\_M.S2 clones from which 3' ESTs had been derived was used as a driver in a hybridization with the NIH\_BMAP\_M.S2 library in the form of single-stranded circles. The remaining single-stranded circles (subtracted library) was purified by hydroxyapatite column chromatography, converted to double-stranded circles and electroporated into DH10B bacteria (Life Technologies) to generate the NIH\_BMAP\_M.S3.3 library. This procedure has been previously described (Bonaldo, Lennon and Soares, Genome Research 6:791-806, 1996).

Research 6:791-806, 1996

NIH\_BMAP\_M.S3.3

TAG-LIB=NIH\_BMAP\_M.S3.3

TAG-TISSUE=basal.ganglia

TAG\_SEQ=GTGAG

BASE COUNT 105 a 87 c 67 g 120 t 1 others

ORIGIN

Query Match 93.6%; Score 23.4; DB 10; Length 380;  
Best Local Similarity 96.0%; Pred. No. 2.8;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 CCATGTGACCATAGGAATGAGAG 25  
|||||  
Db 72 CCATGTGACCATAGGAATGAGAG 48

# RESULT 4

AM124493

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

FEATURES

source

Location/Qualifiers

1. 386

/organism="Mus musculus"

/strain="C57BL/6J"

/db\_xref="taxon:10090"

/clone="UI-M-BH2.1-apo-a-05-0-UI"

/clone\_lib="NIH\_BMAP\_M.S3.1"

/dev\_stage="27-32 days"

/lab\_host="DH10B (Life Technologies)"

/note="Vector: pT73D-Pac (Pharmacia) with a modified

BASE COUNT 106 a 72 c 91 g 117 t

ORIGIN

Query Match 93.6%; Score 23.4; DB 10; Length 386;  
Best Local Similarity 96.0%; Pred. No. 2.8;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 CCATGTGACCATAGGAATGAGAG 25  
|||||  
Db 315 CCATGTGACCATAGGAATGAGAG 339

# RESULT 5

BE980521

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

polylinker. Site 1: Not I; Site 2: Eco RI; The NIH\_BMAP\_M.S3.1 library is a subtracted library of a series, ultimately derived from a mixture of individually tagged, normalized libraries from ten regions of the mouse brain (cerebellum, brain stems, olfactory bulbs, hypothalamus, cortex, amygdala, basal ganglia, pineal gland, striatum, hippocampus) after a series of subtractions to reduce the representation of cDNAs from which ESTs had already been generated. The following process: NIH\_BMAP\_M.S3.1, NIH\_BMAP\_M.S2, NIH\_BMAP\_M.S1. The subtracted library (NIH\_BMAP\_M.S3.1) was constructed as follows: PCR amplified cDNA inserts from NIH\_BMAP\_M.S2 clones from which 3' ESTs had been derived was used as a driver in a hybridization with the NIH\_BMAP\_M.S2 library single-stranded circles (subtracted library). The remaining by hydroxyapatite column chromatography, converted to double-stranded circles and electroporated into DH10B bacteria (Life Technologies) to generate the NIH\_BMAP\_M.S3.1 library. This procedure has been previously described (Bonaldo, Lennon and Soares, Genome Research 6:791-806, 1996).

Research 6:791-806, 1996

NIH\_BMAP\_M.S3.1

TAG-LIB=NIH\_BMAP\_M.S3.1

TAG-TISSUE=amygdala

TAG\_SEQ=GTGAG

limited collaborative arrangements  
Seq primer: M13 Forward  
POLYA-yes.

# FEATURES

Location/Qualifiers  
1. .395

/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UI-M-BG2-bck-b-03-0-UI"  
/clone\_lib="NIH\_BMAP\_MSC\_S1"  
/dev\_stage="27-32 days"  
/lab\_host="DHI0B (Life Technologies)"  
/note="Vector: pT73D-Pac (Pharmacia) with a modified  
polylinker; Site\_1: Not I; Site\_2: Eco RI; The  
NIH\_BMAP\_MSC\_S1 library is a subtracted library derived  
from NIH\_BMAP\_MSC\_N. NIH\_BMAP\_MSC\_N was made from mouse spinal  
cord tissue. For a detailed description of the library  
from which this clone was derived, please visit our web  
site at brainest.eng.uiowa.edu.  
TAG\_LIB=NIH\_BMAP\_MSC\_S1  
TAG\_TISSUE=amygdala  
TAG\_SEQ=GTGAG"

BASE COUNT 108 a 73 c 93 g 121 t

## ORIGIN

Query Match 93.6%; Score 23.4; DB 12; Length 395;  
Best Local Similarity 96.0%; Pred. No. 2.9;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATTTAGGAATGAG 25

Db 315 CCATGTGACCATTTAGGAATGAG 339

## RESULT 6

BF406698

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

BF406698 501 bp mRNA linear EST 28-NOV-2000  
UI-R-BJ2-bpo-b-02-0-UI.s1 UI-R-BJ2 Rattus norvegicus cDNA clone  
UI-R-BJ2-bpo-b-02-0-UI 3', mRNA sequence.  
BF406698  
BF406698.1 GI:11394673  
EST.  
Norway rat.  
Rattus norvegicus  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
Rattus.  
1 (bases 1 to 501)  
Bonaldo,M.F., Lennon,G. and Soares,M.B.  
Normalization and subtraction: two approaches to facilitate gene  
discovery  
Genome Res. 6 (9), 791-806 (1996)  
9704477  
Contact: Soares, MB  
Program for Rat Gene Discovery and Mapping  
University of Iowa  
451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: msoares@blue.weeg.uiowa.edu  
Oligo-dT track not found. Not 1 site shown in beginning of sequence  
is likely internal to the message. cDNA Library Preparation: M.B.  
Soares Lab Clone distribution: clones will be available through  
Research Genetics (www.resgen.com) The following repetitive  
elements were found in this cDNA sequence: 25-96, >B1-F#SINE/Alu  
Seq primer: M13 Forward  
POLYA-No.

## FEATURES

source

Location/Qualifiers  
1. .501  
/organism="Rattus norvegicus"  
/strain="Sprague-Dawley"  
/db\_xref="taxon:10116"  
/clone="UI-R-BJ2-bpo-b-02-0-UI"

BASE COUNT 141 a 100 c 125 g 135 t

## ORIGIN

Query Match 93.6%; Score 23.4; DB 12; Length 501;  
Best Local Similarity 96.0%; Pred. No. 3.2;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATTTAGGAATGAG 25

Db 473 CCATGTGACCATTTAGGAATGAG 497

## RESULT 7

A1481068/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

A1481068 591 bp mRNA linear EST 09-MAR-1999  
vfe2d03.x1 Soares,mammary.gland.NbMWG Mus musculus cDNA clone  
IMAGE:851237 3', similar to FR:008741 008741 HYPXIA-INDUCIBLE  
FACTOR ONE ALPHA.1, mRNA sequence.  
A1481068  
A1481068.1 GI:4374294  
EST.  
house mouse.  
Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 591)  
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
Tumor Gene Index  
Unpublished (1997)  
Contact: Robert Strausberg, Ph.D.  
Email: cgapbs-r@mail.nih.gov  
This clone is available royalty-free through LNL; contact the  
IMAGE Consortium (info@image.llnl.gov) for further information.  
MGI:503389  
This clone was previously sequenced on the 5' end only, this new  
data is from the 3' end  
possible reversed clone; similarity on wrong strand  
High quality sequence stop: 472.

## FEATURES

source

Location/Qualifiers  
1. .591  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="IMAGE:851237"  
/clone\_lib="Soares,mammary.gland.NbMWG"  
/sex="male"  
/tissue\_type="mammary gland"  
/dev\_stage="4 weeks"  
/lab\_host="DHI0B"  
/note="Organ: mammary gland; Vector: pT73D-Pac (Pharmacia  
with a modified polylinker; Site\_1: Not I; Site\_2: Eco  
RI; 1st strand cDNA was primed with a Not I - Oligo(dT)  
primer [5'  
TGTTCACCAATCGAAGTGGAGCGCCGCAATGCTTTTCTTTTCTTTTCTTTT  
T 3']; double-stranded cDNA was ligated to Eco RI  
adaptors (Pharmacia), digested with Not I and cloned into  
the Not I and Eco RI sites of the modified pT733 vector.  
RNA provided by Dr. Minoru Ko, Wayne State Univ. Library  
constructed and normalized by Bento Soares and M.Fatima



BASE COUNT 150 a 143 c 116 g 182 t  
ORIGIN

Query Match 93.6%; Score 23.4; DB 9; Length 591;  
Best Local Similarity 96.0%; Pred. No. 3.4;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATAGGAATGAGAG 25  
|||||  
Db 390 CCATGTGACCATAGGAATGAGAG 366

RESULT 8  
BM834484

LOCUS 684 bp mRNA linear EST 06-MAR-2002  
DEFINITION K-EST0109485 S1LSNU1 Homo sapiens CDNA clone S1LSNU1-63-B11 5',  
BM834484  
BM834484.1 GI:19190893  
ACCESSION  
KEYWORDS  
SOURCE  
ORGANISM

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

Unpublished (2002)  
Contact: Kim YS  
Genome Research Institute of Bioscience & Biotechnology  
52 Deoun-dong Yuseong-gu, Daejeon 305-333, South Korea  
Tel: +82-42-860-4470  
Fax: +82-42-860-4409  
Email: yongsungemail.kr@ib.re.kr  
Plate: 63 row: B column: 11  
High quality sequence stop: 684.  
Location/Qualifiers  
1. 684  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="S1LSNU1-63-B11"  
/clone\_lib="S1LSNU1"  
/sex="M"  
/tissue\_type="Stomach"  
/cell\_line="Lymphoblast-like"  
/lab\_host="Top10F"  
/note="Organ: Stomach; Vector: pME18-FL3; Site: 1: XhoI;  
Site: 2: XhoI; The poly (A)+ RNA was dephosphorylated with  
bacterial alkaline phosphatase (BAP) and then deprotected  
with tobacco acid pyrophosphatase (TAP). The deprotected  
intact mRNA was ligated with DNA-RNA linker including SfiI  
site by treatment of T4 RNA ligase and the first strand  
cDNA was synthesized with Superscript II using SfiI  
oligo-dT primer. After first strand synthesis, RNA was  
degraded by NaOH treatment and cDNA was amplified by PCR  
reaction. The PCR products were digested with SfiI and  
cloned into DraIII-digested pME18S-FL3 vector. The  
obtained cDNA vectors were used for transformation of  
competent cells E. coli Top10F by electroporation method.  
The cDNA libraries constructed by this method are  
full-length enriched cDNA library."

BASE COUNT  
ORIGIN

Query Match 93.6%; Score 23.4; DB 14; Length 684;  
Best Local Similarity 96.0%; Pred. No. 3.6;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATAGGAATGAGAG 25  
|||||  
Db 75 CCATGTGACCATAGGAATGAGAG 99

RESULT 9  
BBI99902

LOCUS 693 bp mRNA linear EST 19-OCT-2001  
DEFINITION BBI99902 RIKEN full-length enriched, 0 day neonate thymus Mus  
musculus CDNA clone A430018A13 3', mRNA sequence.  
BBI99902  
BBI99902.2 GI:16271363  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

Unpublished (2001)  
On Jun 30, 2000 this sequence version replaced gi:8864855.  
Contact: Yoshinobu Hayashizaki  
Laboratory for Genome Exploration Research Group, RIKEN Genomic  
Sciences Center(GSC), Yokohama Institute  
The Institute of Physical and Chemical Research (RIKEN)  
1-7-22 Suenho-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
Tel: 81-45-503-9222  
Fax: 81-45-503-9216  
Email: genome-res@gscc.riken.go.jp,  
URL: http://genome.gsc.riken.go.jp/  
Carninci, P. Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh  
, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.  
Normalization and subtraction of cap-trapper-selected cDNAs to  
prepare full-length cDNA libraries for rapid discovery of new  
genes. Genome Res. 10 (10), 1617-1630 (2000)  
wagi, K., Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,  
Watanabe, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura  
, S., Kawai, T., Okazaki, Y., Muramatsu, M., Inoue, Y., Kita, A. and  
Hayashizaki, Y.  
RIKEN integrated sequence analysis (RISA) system--384-format  
sequencing pipeline with 384 multichannel sequencer. Genome Res.  
10 (11), 1757-1771 (2000)  
Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara  
, Y. and Hayashizaki, Y.  
Computer-based methods for the mouse full-length cDNA  
encyclopedia: real-time sequence clustering for construction of a  
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)  
Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamane, I., Aizawa  
, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and  
Hayashizaki, Y.  
Computational Analysis of Full-length Mouse cDNAs Compared with  
Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)  
Please visit our web site (http://genome.gsc.riken.go.jp/) for  
further details.  
cDNA library was prepared and sequenced in Mouse Genome  
Encyclopedia Project of Genome Exploration Research Group in Riken  
Genomic Sciences Center and Genome Science Laboratory in RIKEN.  
Division of Experimental Animal Research in Riken contributed to  
prepare mouse tissues.

FEATURES  
source

1. 693  
Location/Qualifiers  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="A430018A13"  
/clone\_lib="RIKEN full-length enriched, 0 day neonate  
thymus"

```

/tissue.type="thymus"
/dew_stage="0 day neonate"
/lab_host="DH108"
/notes="Site_1: Sall; Site_2: BamHI. cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN, Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
primed with a primer [5'
GAGGAGAGAGAGATCCAGACGCTCTTTTCTTTTCTTTTNN 3']. cDNA was
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. cDNA went through one round of normalization
to Rot = 10.0 and subtraction to Rot = 459.0. Second
strand cDNA was prepared with the primer adaptor of
sequence [5' GAGGAGAGAGATTCTCCAGTTAATTAATTATATCCCCCCCC
3']. cDNA was cleaved with XhoI and BamHI. Vector: a
modified pluescript KS(+) after bulk excision from Lambda
FIC I."
```

Query Match:	93.6%	Score 23.4	DB 10	Length 693
Best Local Similarity	96.0%	Pred. No. 3.6		
Matches 24	Conservative	0	Mismatches	1
			Indels	0
			Gaps	0
QY	1	CCATGTGACCATAGCAATGAG	25	
Db	414	CCATGTGACCATAGCAATGAG	438	

RESULT 10					
BE731662					
LOCUS	BE731662	711 bp	mRNA	linear	EST 15-SEP-2000
DEFINITION	60156704.8F1 NH_MGC_21 Homo sapiens cDNA clone IMAGE:3842146 5', mRNA sequence.				

VERSION	BE731662.1	GI:10145654
KEYWORDS	EST.	
SOURCE	human.	
ORGANISM	Homo sapiens	

REFERENCE  
1 (Pages 1 to 71)  
AUTHORS NIH-MGC <http://mgc.ni.nih.gov/>.  
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished (1999)  
COMMENT Contact: Robert Strausberg, Ph.D.  
[strausberg@nsl.jhu.edu](mailto:strausberg@nsl.jhu.edu)

CDNA Library Preparation: Ling Hong/Rubin Laboratory  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LNL at: [image.llnl.gov](http://image.llnl.gov)  
 Plate: LINC534 row: 0 column: 11  
 High quality sequence stop: 697.  
 Location/Qualifiers

```

SOURCE
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3842146"
/clone_1ib="NH_MGC_21"
/tissue_type="choriocarcinoma"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: Placenta; Vector: pOTR; Site_1: XhoI;
Site_2: EcoRI; cDNA made by oligo-dT priming.
Directionally cloned into EcoRI/XhoI sites using the
following 5' adaptor: GGCACGAG(G). Size-selected >500bp
for average insert size 1.8kb. Library constructed by
ling Hong in the laboratory of Gerald M. Rubin (University
of California, Berkeley) using ZAP-cDNA synthesis kit

```

BASE COUNT	(Stratagene) and Superscript II RT (Life Technologies),"			
ORIGIN	174 a	166 c	197 g	174 t

Query Match	93.68;	Score 23.4;	DB 12;	Length 711;
Best Local Similarity	96.08;	Pred. No. 3.6;		
Matches 24;	Conservative	0;	Mismatches 1;	Indels 0;
				Gaps 0;

QY 1 CCATGTGCCATTAGGAATGAG 25  
|||||  
Db 646 CCATGTGCCATTAGGAATGAG 670

RESULT 11	734 bp	linear	EST 12-JUN-2001
BE789080		mRNA	
LOCUS			
DEFINITION			
60210490f1 NCI CGAP_Kid14 Mus musculus cDNA clone IMAGE:4223096			
5', mRNA sequence.			

VERSION BF/859080.1 GI:12034110  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus

REFERENCE  
1 (Dases I to 734)  
AUTHORS  
NIH-MGC <http://mgc.ncl.nih.gov/>.  
TITLE  
National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL  
Unpublished (1999)  
COMMENT  
Contact: Robert Strausberg, Ph.D.  
[robertf@mail.nih.gov](mailto:robertf@mail.nih.gov)

Tissue Procurement: Jeffrey E. Green, M.D.  
 CDNA Library Preparation: Life Technologies, Inc.  
 DNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)  
 CDNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LNL at:  
<http://image.llnl.gov>  
 Plate: LAMB810 row: p column: 09  
 High quality sequence stop: 708.

FEATURES	LOCATION/CONTENTS
SOURCE	1. .734

```

/strain="FVB/N"
/db_xref="taxon:10090"
/clone="IMAGE:4223096"
/clone_1b="NCL CGAP Kid14"
/lab_host="PH10B /r1 phage-resistant"
/note="Organ: Kidney; Vector: PCWV-SpORf6; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally. Primer: oligo dF.
Average insert size 1.75 kb. Constructed by Life
Technologies. Note: this is a NCL-CGAP Library."

```

BASE COUNT	AT	T	C	G	
ORIGIN					

Query Match	93.68;	Score 23.4;	DB 12;	Length 734;
Best Local Similarity	96.08;	Pred. No. 3.7;		
Matches 24; Conservative	0;	Mismatches 1;	Indels 0;	Gaps 0

QY 1 CCATGTGACCATTTAGGAATGAGAG 25  
|||||  
Db 306 CCATGTGACCATGAGGAATGAGAG 330

RESULT 12					
BI413382					
LOCUS	6029867331	NCI_CGAP_Lu33	Mus musculus	cDNA clone	IMAGE:5142800 5'
DEFINITION	BI413382		739 bp	mRNA	linear
ACCESSION	BI413382				
VERSION	BI413382.1				
KEYWORDS	EST				
SOURCE	house mouse.				

## ORGANISM

Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 739)  
NIH-MGC <http://imgc.ncl.nih.gov/>.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: [cgabbs-remail.nih.gov](mailto:cgabbs-remail.nih.gov)  
Tissue Procurement: Gilbert Smith, Ph.D.  
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima  
Bonaldo, Ph.D.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: NCI-CGAP clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:  
<http://image.llnl.gov>  
Plate: LLM11352 row: a column: 09  
High quality sequence start: 125  
High quality sequence stop: 726.  
Location/Qualifiers  
1. 739

## FEATURES

source

/organism="Mus musculus"  
/strain="C57BL/6J"  
/db.xref="taxon:10090"  
/clone="IMAGE:5142800"  
/clone.lib="NCI-CGAP\_Lu33"  
/tissue\_type="pooled lung tumors"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: lung; Vector: pRT73D-Pac (Pharmacia) with a  
modified polylinker; Site\_1: NotI; Site\_2: EcoRI; 1st  
lung tumors with a Not I - oligo(dT) primer [5].  
TGTACCACTCTAGAGTGGAGCGCCCTCTCTTTTCTTTTCTTTT 3' ].  
Double-stranded cDNA was ligated to Eco RI adaptors  
(Pharmacia), digested with Not I and cloned into the Not  
I and Eco RI sites of the modified pRT73 vector. Library  
went through one round of normalization, and was  
constructed by Bento Soares and M. Fatima Bonaldo. "

## BASE COUNT

180 a 176 c 210 g 171 t

## Query Match

Best Local Similarity 93.6%; Score 23.4; DB 13; Length 739;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCATGTGACCATTTAGGAATGAGAG 25  
|||||  
Db 681 CCATGTGACCATTTAGGAATGAGAG 705

## RESULT 13

BG772697

LOCUS

## DEFINITION

602720844F1 NIH\_MGC\_97 Homo sapiens CDNA clone IMAGE:4837691 5',  
mRNA sequence.

Accession

## VERSION

BG772697.1 GI:14083350

## KEYWORDS

EST.

SOURCE

human.

## ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 769)  
NIH-MGC <http://imgc.ncl.nih.gov/>.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: [cgabbs-remail.nih.gov](mailto:cgabbs-remail.nih.gov)  
Tissue Procurement: Niklos Palkovits, M.D., Ph.D.  
CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shitaki  
Toshiyuki and Piero Carninci (RIKEN)  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)

## REFERENCE

1

## AUTHORS

TITLE

## JOURNAL

COMMENT

## FEATURES

source

DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:  
<http://image.llnl.gov>  
Plate: LLM10770 row: h column: 12  
High quality sequence stop: 723.  
Location/Qualifiers  
1. 769  
/organism="Homo sapiens"  
/db.xref="taxon:9606"  
/clone="IMAGE:4837691"  
/clone.lib="NIH\_MGC\_97"  
/lab\_host="DH10B"  
/note="Organ: testis; Vector: pBluescript (modified  
pBluescript KS+); Site\_1: BamHI; Site\_2: SalI-XhoI (9cgcg  
size-selected for average insert size 2.2 kb and  
normalized to ROP 5. This is a primary library enriched  
for full-length clones and constructed using the  
cap-trapper method (Carninci, in preparation). Library  
constructed by M. Brownstein (NHGRI), National  
Institutes of Health). Note: this is a NIH-MGC Library. "

## BASE COUNT

243 a 143 c 170 g 213 t

## Query Match

Best Local Similarity 93.6%; Score 23.4; DB 12; Length 769;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CCATGTGACCATTTAGGAATGAGAG 25  
|||||  
Db 434 CCATGTGACCATTTAGGAATGAGAG 458

## RESULT 14

BG110077

LOCUS

## DEFINITION

602279683F1 NIH\_MGC\_86 Homo sapiens CDNA clone IMAGE:4367229 5',  
mRNA sequence.

Accession

## VERSION

BG110077.1 GI:12603583

## KEYWORDS

EST.

SOURCE

human.

## ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 791)  
NIH-MGC <http://imgc.ncl.nih.gov/>.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: [cgabbs-remail.nih.gov](mailto:cgabbs-remail.nih.gov)  
Tissue Procurement: ATCC  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:  
<http://image.llnl.gov>  
Plate: LLM10019 row: e column: 22  
High quality sequence stop: 715.  
Location/Qualifiers  
1. 791

## FEATURES

source

/organism="Homo sapiens"  
/db.xref="taxon:9606"  
/clone="IMAGE:4367229"  
/clone.lib="NIH\_MGC\_86"  
/tissue\_type="osteosarcoma, cell line"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: bone; Vector: pCMV-Sport6; Site\_1: NotI;  
Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
Average insert size 1.533 kb. Library enriched for  
full-length clones and constructed by Life Technologies.

Note: this is a NIH\_MGC Library."

BASE COUNT 250 a 145 c 173 g 223 t

ORIGIN

Query Match 93.6%; Score 23.4; DB 12; Length 791;  
Best Local Similarity 96.0%; Pred. No. 3.8;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATAGGAATGAGAG 25  
|||||

Db 236 CCATGTGACCATAGGAATGAGAG 260

# RESULT 15

AU132294 878 bp mRNA linear EST 01-AUG-2002  
LOCUS AU132294 NT2RP3 Homo sapiens cDNA clone NT2RP3004165 5', mRNA  
DEFINITION sequence.

ACCESSION AU132294  
VERSION AU132294.1 GI:10992648

KEYWORDS

SOURCE

human.

EST.

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE

AUTHORS

Ota,T., Nishikawa,T., Suzuki,Y., Ishii,S., Saito,K., Kawai,Y., Yamamoto,J., Wakamatsu,A., Nakamura,Y., Nagai,T., Sugano,S. and Isogai,T.

TITLE

HRI human cDNA project

Unpublished (2000)

Contact: Takao Isogai

Genomics Laboratory

Helix Research Institute

1532-3 Yana, Kisarazu, Chiba 292-0812, Japan

Tel: 81-438-52-3975

Fax: 81-438-52-3986

Email: genomics@hri.co.jp

HRI human cDNA project; 5'- & 3'-end one pass sequencing; Helix

Research Institute; cDNA library construction; Department of

Virology, Institute of Medical Science, University of Tokyo, and

Helix Research Institute.

FEATURES

Location/Qualifiers

1..878

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="NT2RP3004165"

/clone\_lib="NT2RP3"

/cell\_type="teratocarcinoma"

/cell\_line="NT2"

/note="Vector: pMT18SFL3; mRNA from NT2 neuronal precursor

cells after 2-weeks retinoic acid (RA) induction"

5 others

BASE COUNT 274 a 163 c 188 g 248 t

ORIGIN

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Best Local Similarity 96.0%; Pred. No. 4;

Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 CCATGTGACCATAGGAATGAGAG 25  
|||||

Db 375 CCATGTGACCATAGGAATGAGAG 399

Search completed: March 17, 2003, 13:09:12  
Job time : 888.737 secs

GenCore version 5.1.4\_p5\_4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 09:43:42 ; Search time 220.903 Seconds

(without alignments)  
3161.870 Million cell updates/sec

Title: US-09-836-439-3

Perfect score: 24

Sequence: 1 gcttcttccagagcgccgca 24

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :  
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2: gb\_htg:\*  
3: gb\_in:\*  
4: gb\_om:\*  
5: gb\_ov:\*  
6: gb\_pat:\*  
7: gb\_ph:\*  
8: gb\_pl:\*  
9: gb\_pr:\*  
10: gb\_ro:\*  
11: gb\_sts:\*  
12: gb\_sy:\*  
13: gb\_un:\*  
14: gb\_vl:\*  
15: em\_ba:\*  
16: em\_fun:\*  
17: em\_hum:\*  
18: em\_in:\*  
19: em\_mu:\*  
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22: em\_ov:\*  
23: em\_pat:\*  
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37: em\_htg\_vrt:\*  
38: em\_sy:\*  
39: em\_higo\_hum:\*  
40: em\_higo\_mus:\*  
41: em\_higo\_other:\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	21.4	89.2	175	5	AF249160
2	21.4	89.2	1056	5	SPRY18677
3	21.4	89.2	2577	5	CLRH11
4	21.4	89.2	3016	6	I18746
5	21.4	89.2	5385	9	AB065668
6	21.4	89.2	6953	6	I18747
7	21.4	89.2	6953	7	HS049742
8	21.4	89.2	163297	2	AC023162
9	21.4	89.2	164396	2	HSAC000380
10	21.4	89.2	168551	9	AC080007
11	20.4	85.0	867	5	U97272
12	19.8	82.5	175	5	AF249130
13	19.8	82.5	175	5	AF249133
14	19.8	82.5	175	5	AF249134
15	19.8	82.5	175	5	AF249136
16	19.8	82.5	175	5	AF249137
17	19.8	82.5	175	5	AF249138
18	19.8	82.5	175	5	AF249139
19	19.8	82.5	175	5	AF249140
20	19.8	82.5	175	5	AF249141
21	19.8	82.5	175	5	AF249142
22	19.8	82.5	175	5	AF249143
23	19.8	82.5	175	5	AF249144
24	19.8	82.5	175	5	AF249146
25	19.8	82.5	175	5	AF249147
26	19.8	82.5	175	5	AF249148
27	19.8	82.5	175	5	AF249149
28	19.8	82.5	175	5	AF249150
29	19.8	82.5	175	5	AF249151
30	19.8	82.5	175	5	AF249152
31	19.8	82.5	175	5	AF249154
32	19.8	82.5	175	5	AF249156
33	19.8	82.5	175	5	AF249157
34	19.8	82.5	175	5	AF249158
35	19.8	82.5	726	5	AF369050
36	19.8	82.5	858	5	AF137208
37	19.8	82.5	864	5	AF137206
38	19.8	82.5	867	5	U97265
39	19.8	82.5	867	5	U97266
40	19.8	82.5	867	5	U97274
41	19.8	82.5	1047	4	AF055319
42	19.8	82.5	1053	5	AF021242
43	19.8	82.5	1150	4	AF008847
44	19.8	82.5	1300	10	AF309568
45	19.8	82.5	1389	5	PMY18679

# ALIGNMENTS

RESULT 1  
AF249160 175 bp DNA linear VRT 17-JAN-2001  
LOCUS Philautus charius rhodopsin gene, exon 4 and partial cds.  
DEFINITION  
ACCESSION AF249160  
VERSION AF249160.1 GI:12247249  
KEYWORDS  
SOURCE  
ORGANISM  
Philautes charius.  
Philautes charius  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Rhacophoridae;  
Philautes.  
REFERENCE  
AUTHORS 1 (bases 1 to 175)  
TITLE Bossuyt, F. and Minkovitch, M.C.  
Convergent Adaptive Radiations in Madagascan and Asian Rain Forest Frogs

Reveal Co-variation between Larval and Adult Traits  
 Unpublished  
 2 (bases 1 to 175)  
 Bossuyt, F. and Milinkovitch, M.C.  
 TITLE Direct Submission  
 Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of  
 Molecular Biology and Medicine, rue Jenner and Brachet 12,  
 Gosselies B-6041, Belgium

FEATURES  
 source Location/Qualifiers

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BASE COUNT 32 a 55 c 36 g 52 t  
 ORIGIN

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 Best Local Similarity 95.7%; Pred. No. 5.5;  
 Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTTGCCAAGAGCGCCG 23  
 Db 140 GCTTCTTTGCCAAGAGCTCCG 162

RESULT 2

SPY18677 1056 bp mRNA linear VRT 26-MAY-1999  
 LOCUS Sardinia pilchardus mRNA for opsin.  
 DEFINITION Y18677  
 ACCESSION Y18677.1 GI:4210840  
 KEYWORDS opsin.  
 SOURCE Sardinia pilchardus.  
 ORGANISM Sardinia pilchardus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Actinopterygii; Neopterygii; Teleostei; Clupeomorpha; Clupeidae;  
 Sardinia.

REFERENCE 1 (bases 1 to 1056)  
 ARCHER, S.N. and HIRANO, J.  
 TITLE Comparative analysis of opsins in Mediterranean coastal fish  
 JOURNAL Unpublished  
 AUTHORS 2 (bases 1 to 1056)  
 TITLE Archer, S.N.  
 JOURNAL Direct Submission  
 Submitted (25-JAN-1999) S.N. Archer, International Marine Centre,  
 Localita sa Mardini, 09072 Torregrande, Oristano, ITALY

FEATURES  
 source Location/Qualifiers

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 FVFGMGNIGCFPATLGGELALSLIVLSERNIVYCKPISNRFGENAHVAVAS  
 WPMACAVPPLVGNRIIPGMOCSCGIDYITRAEGNNSEFVIMVAFHFTCLPLI  
 ITFCYGRIVCTVKAADQOSETTORAEREVTVIIMFAVLAICWVPAVASVAMVIF

THGSEFGPVEMTIPAFPAKSSAVNPVYIICLNKQFRHCHMTITLCCGRNPPEEGRS  
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BASE COUNT 186 a 368 c 269 g 233 t  
 ORIGIN

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 Best Local Similarity 95.7%; Pred. No. 6.6;  
 Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTTGCCAAGAGCGCCG 23  
 Db 874 GCTTCTTTGCCAAGAGCTCCG 896

RESULT 3

CLRH11 2577 bp DNA linear VRT 17-DEC-1999  
 LOCUS Columbia livia rh1 opsin (rh1) gene, exons 1 through 4.  
 DEFINITION AF149230  
 ACCESSION AF149230  
 VERSION AF149230.1 GI:4887218  
 KEYWORDS  
 SEGMENT  
 SOURCE 1 of 2  
 ORGANISM Columbia livia.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Archosauria; Aves; Neognathae; Columbiformes; Columbidae; Columba.

REFERENCE 1 (bases 1 to 2577)  
 ARCHER, S.N. and HIRANO, J.  
 TITLE Genetic analyses of visual pigments of the pigeon (Columba livia)  
 JOURNAL Genetics 153 (4), 1839-1850 (1999)  
 MEDLINE 20050679  
 PUBMED 10581289

REFERENCE 2 (bases 1 to 2577)  
 ARCHER, S.N. and HIRANO, J.  
 TITLE Direct Submission  
 Submitted (10-MAY-1999) Department of Biology, Syracuse University,  
 130 College Place, Syracuse, NY 13244, USA  
 JOURNAL Location/Qualifiers

FEATURES  
 source Location/Qualifiers

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 1561..1726  
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 ORIGIN

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 Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTTGCCAAGAGCGCCG 23  
 Db 2145 GCTTCTTTGCCAAGAGCTCCG 2167

RESULT 4  
 LOCUS 118746 3016 bp DNA linear PAT 07-OCT-1996  
 DEFINITION Sequence 1 from patent US 5498521.

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ACCESSION 118746
VERSION 118746.1 GI:1599101
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE Unclassified.
AUTHORS 1 (bases 1 to 3016)
TITLE Dryja,T.P. and Berson,E.L.
JOURNAL Diagnosis of hereditary retinal degenerative diseases
FEATURES
BASE COUNT 689 a 863 c 753 g 711 t
ORIGIN
Query Match
Best Local Similarity 89.2%; Score 21.4; DB 6; Length 3016;
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GCTTCTTTGCCAAGAGCGCCGC 23
Db 1168 GCGTCTTTGCCAAGAGCGCCGC 1190

RESULT 5
LOCUS AB065668
DEFINITION Homo sapiens gene for seven transmembrane helix receptor, complete
ACCESSION AB065668
VERSION AB065668.1 GI:21928610
KEYWORDS
SOURCE
ORGANISM Homo sapiens (Isolate:CBRC7M_231) DNA.
REFERENCE
AUTHORS Suwa,M., Sato,T., Okouchi,I., Arita,M., Putani,K., Matsunoto,S.,
TITLE Tsubumi,S., Aburatani,H., Asai,K. and Akiyama,Y.
JOURNAL Genome-wide discovery and analysis of human seven transmembrane
REFERENCE helix receptor genes
AUTHORS Unpublished
JOURNAL 2 (bases 1 to 5385)
Suwa,M.
COMMENT Direct Submission
Submitted (11-JUL-2001) Makiko Suwa, Computational Biology Research
Center (CBRC), National Institute of Advanced Industrial Science
and Technology (AIST); 2-41-6 Aomi Koto-ku, Tokyo 135-0064, Japan
(E-mail:m-suwa@aist.go.jp, URL:http://www.cbrc.jp/,
Tel:81-3-3599-8080, Fax:81-3-3599-8081)
This sequence is a seven transmembrane helix receptor candidate
predicted from the whole human genome sequences using our automated
finding(GeneDecoder), sequence search, motif-domain assignment and
transmembrane helix prediction.
And the sequence is submitted by the collaborative project between
[Computational Biology Research Center (CBRC), National Institute
of Advanced Industrial Science and Technology (AIST)] and [Genome
Science Division, Research Center for Advanced Science and
Technology (RCAST), University of Tokyo].
FEATURES
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/db_xref="taxon:9606"
/chromosome="3"
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FLIVIGPINFILTYVQHKRLPLNTILNLAVADLFVWLGFTSTLYSLNGY
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WVALACAPPLAGMSRYIPGILQSCGIDYVTLKPEVNNEFVYVFWVHTPIPI
IFECYGLVFTPEKDAADQDSATQKAEKEVPMVITVIAFLICVPPVSAFVIF
THQSNFGPIEMTIPAFEFKSAALYIVYIIMNKKPRNMLTTICGKNPLGDDEAS
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BASE COUNT 1132 a 1618 c 1407 g 1228 t
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Best Local Similarity 89.2%; Score 21.4; DB 9; Length 5385;
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GCTTCTTTGCCAAGAGCGCCGC 23
Db 4177 GCGTCTTTGCCAAGAGCGCCGC 4199

RESULT 6
LOCUS 118747
DEFINITION Sequence 2 from patent US 5498521.
ACCESSION 118747
VERSION 118747.1 GI:1599102
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE Unclassified.
AUTHORS 1 (bases 1 to 6953)
TITLE Dryja,T.P. and Berson,E.L.
JOURNAL Diagnosis of hereditary retinal degenerative diseases
FEATURES
BASE COUNT 1523 a 2022 c 1797 g 1611 t
ORIGIN
Query Match
Best Local Similarity 89.2%; Score 21.4; DB 6; Length 6953;
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GCTTCTTTGCCAAGAGCGCCGC 23
Db 4272 GCGTCTTTGCCAAGAGCGCCGC 4294

RESULT 7
LOCUS HS049742
DEFINITION Human rhodopsin gene, complete cds.
ACCESSION U49742
VERSION U49742.1 GI:1236136
KEYWORDS opsin; rhodopsin.
SOURCE
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
TITLE 1 (bases 1 to 6953)
Nathans,J. and Hogness,D.S.
JOURNAL Isolation and nucleotide sequence of the gene encoding human
rhodopsin
Proc. Natl. Acad. Sci. U.S.A. 81 (15), 4851-4855 (1984)
PUBMED 6589631
REFERENCE
AUTHORS 2 (bases 1 to 6953)
Nathans,J.
JOURNAL Direct Submission
Submitted (22-FEB-1996) Jeremy Nathans, Molecular Biology and
Genetics, Johns Hopkins Medical School, 725 N. Wolfe Street,
Baltimore, MD 21205, USA
On Sep 3, 1996 this sequence version replaced gi:189393.

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SUBMIT	13
P249133	
CDS	
DEFINITION	175 bp DNA linear VRT 17-JAN-2001
ACCSSION	M86181 madagascariensis rhodopsin gene, exon 4 and partial cds.
VERSION	AZ249133
TWORDS	AZ249133.1 GI:12247195
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SOURCE  
ORGANISM  
Mantella madagascariensis.  
Mantella madagascariensis  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Rhacophoridae;  
Mantella.

REFERENCE  
AUTHORS  
Bossuyt, F. and Milinkovitch, M.C.  
TITLE  
Reveal Co-variation between larval and Adult Traits

JOURNAL  
2 (bases 1 to 175)  
Bossuyt, F. and Milinkovitch, M.C.  
REFERENCE  
AUTHORS  
Bossuyt, F. and Milinkovitch, M.C.  
TITLE  
Direct Submission  
Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of  
Molecular Biology and Medicine, rue Jeener and Brachet 12,  
Gosselies B-6041, Belgium

FEATURES  
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33 a 53 c 36 g 52 t 1 others

BASE COUNT  
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Best Local Similarity 91.3%; Pred. No. 37;  
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GCTTCTTTGCCAAGAGCGCCG 23  
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Db 140 GCTTCTTTGCCAAGAGCTCTGC 162

RESULT 14  
AF249134 175 bp DNA linear VRT 17-JAN-2001  
LOCUS  
Mantidactylus cf. ulcerosus rhodopsin gene, exon 4 and partial cds.  
ACCESSION  
AF249134  
VERSION  
AF249134.1 GI:12247197  
KEYWORDS  
Mantidactylus cf. ulcerosus.  
SOURCE  
Mantidactylus cf. ulcerosus.  
ORGANISM  
Mantidactylus cf. ulcerosus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Rhacophoridae;  
Mantidactylus.

REFERENCE  
AUTHORS  
Bossuyt, F. and Milinkovitch, M.C.  
TITLE  
Convergent Adaptive Radiations in Madagascan and Asian Ranid Frogs  
Reveal Co-variation between larval and Adult Traits

JOURNAL  
2 (bases 1 to 175)  
Bossuyt, F. and Milinkovitch, M.C.  
REFERENCE  
AUTHORS  
Bossuyt, F. and Milinkovitch, M.C.  
TITLE  
Direct Submission  
Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of  
Molecular Biology and Medicine, rue Jeener and Brachet 12,  
Gosselies B-6041, Belgium

FEATURES  
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BASE COUNT  
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ORIGIN  
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Best Local Similarity 91.3%; Pred. No. 37;  
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GCTTCTTTGCCAAGAGCGCCG 23  
|||||  
Db 140 GCTTCTTTGCCAAGAGCTCTGC 162

RESULT 15  
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LOCUS  
Boophis xerophilus rhodopsin gene, exon 4 and partial cds.  
ACCESSION  
AF249136  
VERSION  
AF249136.1 GI:12247201  
KEYWORDS  
Boophis xerophilus.  
SOURCE  
Boophis xerophilus.  
ORGANISM  
Boophis xerophilus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Rhacophoridae;  
Boophis.

REFERENCE  
AUTHORS  
Bossuyt, F. and Milinkovitch, M.C.  
TITLE  
Convergent Adaptive Radiations in Madagascan and Asian Ranid Frogs  
Reveal Co-variation between larval and Adult Traits

JOURNAL  
2 (bases 1 to 175)  
Bossuyt, F. and Milinkovitch, M.C.  
REFERENCE  
AUTHORS  
Bossuyt, F. and Milinkovitch, M.C.  
TITLE  
Direct Submission  
Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of  
Molecular Biology and Medicine, rue Jeener and Brachet 12,  
Gosselies B-6041, Belgium

FEATURES  
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exon  
34 a 55 c 34 g 52 t

BASE COUNT  
34 a 55 c 34 g 52 t

ORIGIN  
Query Match 82.5%; Score 19.8; DB 5; Length 175;  
Best Local Similarity 91.3%; Pred. No. 37;  
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GCTTCTTTGCCAAGAGCGCCG 23  
|||||  
Db 140 GCTTCTTTGCCAAGAGCTCTGC 162

Search completed: March 17, 2003, 11:25:45  
Job time : 313.903 secs



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FT misc_binding complement (355..369)
FT /tag= f
FT /note= "Binds probes AAT17119 (mutant) and AAT17120
FT (normal)"
FT mutation 362
FT /tag= g
FT /note= "Substitution with A in mutant sequence"
FT primer_bind 362..381
FT /tag= h
FT /note= "Binds primers 485 (AAT17122) (normal) and 502
FT (mutant)"
FT US5498521-A.
FT 12-MAR-1996.
FT 24-JAN-1990; 90US-0469215.
FT 11-MAR-1993; 93US-0033081.
FT 24-JAN-1990; 90US-0469215.
FT 11-DEC-1991; 91US-0805123.
FT (HARD ) HARVARD COLLEGE.
FT Berson EL, Dryja TP;
FT WPI; 1996-159684/16.
FT P-PSDB; AAR93116.
FT Diagnosis of hereditary retinal degenerative diseases e.g. retinitis
FT pigmentosa, - caused by a human photoreceptor protein mutation, by
FT detection of the mutation by PCR amplification or hybridisation
FT methods
FT Example 1; Column 19-24; 71pp; English.
FT
FT This sequence encodes human rhodopsin, and is shown without
FT introns. The full sequence, with introns, is shown in AAT17116.
FT Substitution of histidine for the normal nonpolar amino acid
FT proline at position 23, by substitution of C with A in codon-23,
FT results in a dysfunctional or absent molecule, affecting rod
FT function, and is linked with autosomal dominant retinitis
FT pigmentosa. Probes AAT17117 and AAT17119 bind to the C-to-A
FT transversion mutation sequence, and probes AAT17118 and AAT17120 bind
FT to the corresponding normal sequence. Primers 485 (AAT17122) and 502
FT (AAT17123) may be used along with primer 348 (AAT17121) to amplify
FT mutant and normal sequences, respectively, by PCR. Mutations in the
FT retinal degeneration slow protein and retinal rod
FT cGMP-phosphodiesterase genes are also implicated in retinitis
FT pigmentosa. Detection of any of these mutations in a foetus or
FT patient may be used in diagnosis.
FT
FT Sequence 3016 BP; 689 A; 863 C; 753 G; 711 T; 0 other;
FT
FT Query Match 89.2%; Score 21.4; DB 17; Length 3016;
FT Best Local Similarity 95.7%; Pred. No. 1.7;
FT Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
FT
FT 1 GCTTCTTTGGCCAGAGCGCCG 23
FT 1168 GCGTCTTTGGCCAGAGCGCCG 1190
FT
FT RESULT 2
FT AAT17116
FT ID AAT17116 standard; DNA: 6953 BP.
FT
FT AAT17116;
FT
FT 06-JUL-1996 (first entry)
FT
FT Rhodopsin gene.
FT
FT

```

```

KW Human: rhodopsin; transversion; mutation; retinitis pigmentosa;
KW intron; probe; primer; hybridisation; polymerase chain reaction; PCR;
KW eye; rod; retina; diagnostic; prenatal diagnosis; photoreceptor; ds.
OS Homo sapiens.
XX Location/Qualifiers
XX Key 200..294
XX 5'UTR /tag= a
XX 202..294
XX 5'UTR /tag= b
XX /note= "Alternative 5'-UTR"
XX primer_bind complement (231..250)
XX /tag= c
XX /note= "Binds primer 348 (AAT17121)"
XX 295..655
XX exon /tag= d
XX /number= 1
XX misc_binding complement (354..372)
XX /tag= e
XX /note= "Binds probes AAT17117 (mutant) and AAT17118
XX complement (355..369)
XX /tag= f
XX /note= "Binds probes AAT17119 (mutant) and AAT17120
XX (normal)"
XX mutation 362
XX /tag= g
XX /note= "Substitution with A in mutant sequence"
XX 362..381
XX primer_bind /tag= h
XX /note= "Binds primers 485 (AAT17122) (normal) and 502
XX (mutant)"
XX intron 666..2438
XX /tag= i
XX /number= 1
XX exon 2439..2607
XX /tag= j
XX /number= 2
XX intron 2608..3812
XX /tag= k
XX /number= 2
XX exon 3813..3978
XX /tag= l
XX /number= 3
XX intron 3979..4094
XX /tag= m
XX /number= 3
XX exon 4095..4334
XX /tag= n
XX /number= 4
XX intron 4335..5167
XX /tag= o
XX /number= 4
XX exon 5168..5278
XX /tag= p
XX /number= 5
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FT US5498521-A.
FT 12-MAR-1996.
FT 24-JAN-1990; 90US-0469215.
FT 11-MAR-1993; 93US-0033081.
FT 24-JAN-1990; 90US-0469215.
FT 11-DEC-1991; 91US-0805123.
FT (HARD ) HARVARD COLLEGE.
FT Berson EL, Dryja TP;
FT WPI; 1996-159684/16.

```

DR P-PSDB; AAR93116.

XX  
PT Diagnosis of hereditary retinal degenerative diseases e.g. retinitis  
PT pigmentosa, - caused by a human photoreceptor protein mutation, by  
PT detection of the mutation by PCR amplification or hybridisation  
PT methods

XX Example 1: Column 23-30; 71pp; English.

XX  
CC This sequence encodes human rhodopsin, and is shown in full with  
CC introns. The corresponding sequence without introns is shown in  
CC AAT17116. Substitution of histidine for the normal nonpolar amino  
CC acid proline at position 23, by substitution of C with A in  
CC codon 23, results in a dysfunctional or absent molecule, affecting  
CC rod function, and is linked with autosomal dominant retinitis  
CC pigmentosa. Probes AAT17117 and AAT17119 bind to the C-to-A  
CC transversion mutation sequence, and probes AAT17118 and AAT17120 bind  
CC to the corresponding normal sequence. Primers 485 (AAT17122) and 502  
CC (AAT17123) may be used along with primer 348 (AAT17121) to amplify  
CC mutant and normal sequences, respectively, by PCR. Mutations in the  
CC cGMP-phosphodiesterase gene are also implicated in retinitis  
CC pigmentosa. Detection of any of these mutations in a foetus or  
CC patient may be used in diagnosis.

XX Sequence 6953 BP; 1523 A; 2022 C; 1797 G; 1611 T; 0 other;

Query Match 89.2%; Score 21.4; DB 17; Length 6953;  
Best Local Similarity 95.7%; Pred. No. 2;

Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTTCTTGGCCAGAGCGCCGC 23  
II |||||||

Db 4272 GCGTCTTGGCCAGAGCGCCGC 4294

#### RESULT 3

AA043543

ID AA043543 standard; cDNA; 8641 BP.

XX AA043543;

DT 08-SEP-1999 (first entry)

DE Stealth virus nucleic acid clone, SEQ ID NO: 36.

XX Stealth virus; detection; diagnosis; infection; ss.

OS Stealth virus.

XX WO934019-A1.

PD 08-JUL-1999.

PF 30-DEC-1998; 98WO-US27744.

PR 30-DEC-1997; 97US-0001184.

XX (MARTIN) MARTIN W J.

PA (MARTIN) MARTIN W J.

PI Martin WJ;

DR WPI; 1999-405521/34.

PT Novel strains of stealth virus

XX Novel strains of stealth virus

PS Claim 19; Page 76-79; 95pp; English.

XX  
CC This sequence represents a Stealth virus nucleic acid clone. The  
CC invention relates to a method of detecting and characterising a stealth  
CC virus by reacting a sample suspected of containing a stealth virus with a  
CC probe from a known stealth virus and sequencing the resultant isolated  
CC nucleotide. The method comprises the steps of: (a) isolating DNA or RNA

CC from a sample suspected of containing a stealth virus, e.g. a culture of  
CC cells showing a viral cytopathic effect; (b) testing the reactivity of  
CC the isolated DNA or RNA with a molecular probe that contains at least 18  
CC or more contiguous nucleotides identical to sequence previously  
CC identified from a stealth virus; and, optionally (c) sequencing the  
CC isolated DNA or RNA molecules that react with the probe. The method is  
CC used to detect stealth virus in a biological product, food or in the  
CC environment. The method is also used to evaluate agents for their  
CC inhibitory or stimulatory effects on stealth virus replication and to  
CC determine capacity of the virus to recombine with and potentially alter  
CC the nucleic acid sequences of a cell or bacterium.

XX Sequence 8641 BP; 2101 A; 2031 C; 2018 G; 2476 T; 15 other;

Query Match 78.3%; Score 18.8; DB 20; Length 8641;  
Best Local Similarity 90.9%; Pred. No. 37;

Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 CTTTCTTGGCCAGAGCGCCGC 23  
II |||||||

Db 8149 CTTTCTTGGCCAGAGCGCCGC 8170

#### RESULT 4

AA043543

ID AA043543 standard; cDNA; 3129 BP.

XX AA043543;

DT 11-NOV-1993 (first entry)

DE Rhodopsin gene.

XX Rhodopsin gene.

XX Human; rhodopsin; mutant; retinal degeneration; primer; probe;

XX hereditary; ss.

OS Homo sapiens.

XX Key

FT prim\_transcript 200..1341

FT CDS 295..1341

FT /tag= a

FT /tag= b

XX WO9312134-A.

PD 24-JUN-1993.

PF 08-DEC-1992; 92WO-US10536.

PR 11-DEC-1991; 91US-0805123.

XX (HARD ) HARVARD COLLEGE.

PA Berson EL, Dryja TP;

PI Berson EL, Dryja TP;

DR WPI; 1993-214088/26.

PT P-PSDB; AAR8483.

XX Probe or primer conty. sequence of human retinal degeneration

XX slow protein mutant - used to diagnose hereditary retinal

XX degenerative diseases

PS Disclosure; Fig 1; 56pp; English.

XX  
CC The sequence given represents the human rhodopsin cDNA. Mutant  
CC versions of this sequence encode proteins which cause retinal  
CC degeneration. These sequences may be identified using primers/  
CC probes described in the invention (see also AA043545-48) and may be  
CC used to diagnose hereditary retinal degeneration. This sequence is  
CC the closest approximation to the gene sequence as the sequence given  
CC in the specification is not printed clearly.

XX AC AAS41923



XX 17-DEC-2001 (first entry)  
XX  
XX Genomic sequence #239 encoding novel human enzyme polypeptide.  
DE  
XX  
XX Human; oxidoreductase enzyme; transferase; hydrolase; lyase; isomerase;  
KW lysase; hyperproliferative disorder; immunodeficiency disorder;  
KW autoimmune disorder; neurological disorder; metabolic disorder;  
KW inflammatory disorder; cardiovascular disorder; reproductive disorder;  
KW blood-related disorder; infectious disorder; gene therapy; cytostatic;  
KW anti arthritic; nephrotoxic; anticoagulant; ds.  
XX  
OS Homo sapiens.  
XX  
XX MO20015301-A2.  
XX  
XX 02-AUG-2001.  
XX  
XX 17-JAN-2001; 2001MO-US01339.  
XX  
XX 31-JAN-2000; 2000US-0179065.  
XX 04-FEB-2000; 2000US-0180628.  
XX 24-FEB-2000; 2000US-0184664.  
XX 02-MAR-2000; 2000US-0186350.  
XX 15-MAR-2000; 2000US-0189874.  
XX 17-MAR-2000; 2000US-0190076.  
XX 18-APR-2000; 2000US-0198123.  
XX 19-MAY-2000; 2000US-0205415.  
XX 07-JUN-2000; 2000US-0209467.  
XX 28-JUN-2000; 2000US-0214886.  
XX 30-JUN-2000; 2000US-0215135.  
XX 07-JUL-2000; 2000US-0216647.  
XX 11-JUL-2000; 2000US-0216880.  
XX 11-JUL-2000; 2000US-0217487.  
XX 14-JUL-2000; 2000US-0218290.  
XX 26-JUL-2000; 2000US-0220963.  
XX 26-JUL-2000; 2000US-0220964.  
XX 14-AUG-2000; 2000US-0224518.  
XX 14-AUG-2000; 2000US-0224519.  
XX 14-AUG-2000; 2000US-0225213.  
XX 14-AUG-2000; 2000US-0225214.  
XX 14-AUG-2000; 2000US-0225266.  
XX 14-AUG-2000; 2000US-0225267.  
XX 14-AUG-2000; 2000US-0225268.  
XX 14-AUG-2000; 2000US-0225270.  
XX 14-AUG-2000; 2000US-0225271.  
XX 14-AUG-2000; 2000US-0225757.  
XX 14-AUG-2000; 2000US-0225758.  
XX 14-AUG-2000; 2000US-0225759.  
XX 18-AUG-2000; 2000US-0226279.  
XX 22-AUG-2000; 2000US-0226681.  
XX 22-AUG-2000; 2000US-0226688.  
XX 22-AUG-2000; 2000US-0227182.  
XX 23-AUG-2000; 2000US-0227189.  
XX 30-AUG-2000; 2000US-0228294.  
XX 01-SEP-2000; 2000US-0229287.  
XX 01-SEP-2000; 2000US-0229343.  
XX 01-SEP-2000; 2000US-0229344.  
XX 01-SEP-2000; 2000US-0229345.  
XX 05-SEP-2000; 2000US-0229509.  
XX 05-SEP-2000; 2000US-0229513.  
XX 06-SEP-2000; 2000US-0230437.  
XX 06-SEP-2000; 2000US-0230438.  
XX 08-SEP-2000; 2000US-0231242.  
XX 08-SEP-2000; 2000US-0231243.  
XX 08-SEP-2000; 2000US-0231244.  
XX 08-SEP-2000; 2000US-0231413.  
XX 08-SEP-2000; 2000US-0231414.  
XX 08-SEP-2000; 2000US-0232080.  
XX 08-SEP-2000; 2000US-0232081.  
XX 12-SEP-2000; 2000US-0231968.  
XX 14-SEP-2000; 2000US-0232397.

PR 14-SEP-2000; 2000US-0232398.  
PR 14-SEP-2000; 2000US-0232399.  
PR 14-SEP-2000; 2000US-0232400.  
PR 14-SEP-2000; 2000US-0232401.  
PR 14-SEP-2000; 2000US-0232403.  
PR 14-SEP-2000; 2000US-0233063.  
PR 14-SEP-2000; 2000US-0233064.  
PR 21-SEP-2000; 2000US-0233065.  
PR 21-SEP-2000; 2000US-0234223.  
PR 25-SEP-2000; 2000US-0234274.  
PR 25-SEP-2000; 2000US-0234997.  
PR 25-SEP-2000; 2000US-0234998.  
PR 26-SEP-2000; 2000US-0235484.  
PR 27-SEP-2000; 2000US-0235834.  
PR 27-SEP-2000; 2000US-0235836.  
PR 29-SEP-2000; 2000US-0236327.  
PR 29-SEP-2000; 2000US-0236367.  
PR 29-SEP-2000; 2000US-0236368.  
PR 29-SEP-2000; 2000US-0236369.  
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PR 02-OCT-2000; 2000US-0236802.  
PR 02-OCT-2000; 2000US-0237037.  
PR 02-OCT-2000; 2000US-0237038.  
PR 02-OCT-2000; 2000US-0237039.  
PR 13-OCT-2000; 2000US-0237040.  
PR 13-OCT-2000; 2000US-0239335.  
PR 20-OCT-2000; 2000US-0240960.  
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PR 20-OCT-2000; 2000US-0241787.  
PR 20-OCT-2000; 2000US-0241808.  
PR 20-OCT-2000; 2000US-0241809.  
PR 01-NOV-2000; 2000US-0244826.  
PR 08-NOV-2000; 2000US-0244874.  
PR 08-NOV-2000; 2000US-0244875.  
PR 08-NOV-2000; 2000US-0246475.  
PR 08-NOV-2000; 2000US-0246476.  
PR 08-NOV-2000; 2000US-0246477.  
PR 08-NOV-2000; 2000US-0246478.  
PR 08-NOV-2000; 2000US-0246523.  
PR 08-NOV-2000; 2000US-0246524.  
PR 08-NOV-2000; 2000US-0246525.  
PR 08-NOV-2000; 2000US-0246526.  
PR 08-NOV-2000; 2000US-0246527.  
PR 08-NOV-2000; 2000US-0246528.  
PR 08-NOV-2000; 2000US-0246529.  
PR 08-NOV-2000; 2000US-0246532.  
PR 08-NOV-2000; 2000US-0246609.  
PR 08-NOV-2000; 2000US-0246610.  
PR 08-NOV-2000; 2000US-0246611.  
PR 08-NOV-2000; 2000US-0246612.  
PR 17-NOV-2000; 2000US-0249207.  
PR 17-NOV-2000; 2000US-0249208.  
PR 17-NOV-2000; 2000US-0249209.  
PR 17-NOV-2000; 2000US-0249210.  
PR 17-NOV-2000; 2000US-0249211.  
PR 17-NOV-2000; 2000US-0249212.  
PR 17-NOV-2000; 2000US-0249213.  
PR 17-NOV-2000; 2000US-0249214.  
PR 17-NOV-2000; 2000US-0249215.  
PR 17-NOV-2000; 2000US-0249216.  
PR 17-NOV-2000; 2000US-0249217.  
PR 17-NOV-2000; 2000US-0249218.  
PR 17-NOV-2000; 2000US-0249244.  
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PR 17-NOV-2000; 2000US-0249264.  
PR 17-NOV-2000; 2000US-0249265.  
PR 17-NOV-2000; 2000US-0249297.  
PR 17-NOV-2000; 2000US-0249299.  
PR 17-NOV-2000; 2000US-0249300.  
PR 01-DEC-2000; 2000US-0250160.  
PR 01-DEC-2000; 2000US-0250391.  
PR 05-DEC-2000; 2000US-0251030.

PR 05-DEC-2000; 2000US-0251988.  
PR 05-DEC-2000; 2000US-0256719.  
PR 06-DEC-2000; 2000US-0251479.  
PR 08-DEC-2000; 2000US-0251856.  
PR 08-DEC-2000; 2000US-0251868.  
PR 08-DEC-2000; 2000US-0251869.  
PR 08-DEC-2000; 2000US-0251989.  
PR 08-DEC-2000; 2000US-0251990.  
PR 11-DEC-2000; 2000US-0254097.  
PR 05-JAN-2001; 2001US-0259678.  
XX  
XX (HUMA-) HUMAN GENOME SCI INC.  
XX  
XX Rosen CA, Barash SC, Ruben SM;  
XX  
XX WPI; 2001-465566/50.  
XX  
XX Novel polypeptides and polynucleotides useful for diagnosing,  
XX preventing, treating neural, immune system, muscular, reproductive,  
XX pulmonary, cardiovascular, renal, proliferative disorders and cancerous  
XX diseases  
XX  
XX Disclosure; SEQ ID NO 2049; 1180pp; English.  
XX  
XX The present invention relates to the isolation of novel human enzyme  
XX polypeptides (AAU22915-AAU23814), and the cDNA and genomic sequences  
XX encoding them. The enzyme polypeptides of the invention may comprise the  
XX functional classes of oxidoreductases, transferases, hydrolases, lyases,  
XX isomerases or ligases. The sequences of the invention are useful in the  
XX diagnosis, treatment, prevention and/or prognosis of a wide range of  
XX disorders including hyperproliferative disorders (e.g. cancer),  
XX immunodeficiency disorders (e.g. AIDS) autoimmune disorders  
XX (e.g. arthritis), neurological disorders (e.g. Alzheimer's disease),  
XX metabolic disorders (e.g. phenylketonuria), inflammatory disorders  
XX (e.g. asthma), cardiovascular disorders (e.g. atherosclerosis),  
XX blood-related disorders (e.g. haemophilia), reproductive disorders  
XX (e.g. infertility) and infectious disorders (e.g. influenza). The  
XX polynucleotides of the invention can also be used in gene therapy.  
XX AAU1665-AAU2192 represent DNA sequences encoding for the novel human  
XX enzyme polypeptides of the invention.  
XX Note: The sequence data for this patent did not form part of the printed  
XX specification, but was obtained in electronic format directly from WIPO  
XX at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
XX Sequence 2209 BP; 629 A; 412 C; 442 G; 726 T; 0 other;  
XX  
XX Query Match 72.5%; Score 17.4; DB 22; Length 2209;  
XX Best Local Similarity 94.7%; Pred. No. 1.4e+02;  
XX Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
XX  
XX 1 GCTTCTTGCCAGAGCG 19  
XX ||||||||||||||||  
XX Db 843 GCTTCTTGCCAGAGCG 861  
XX  
XX RESULT 8  
XX AAU90036/c  
XX ID AAU90036 standard; DNA; 2601 BP.  
XX  
XX AAU90036;  
XX  
XX 17-SEP-1999 (first entry)  
XX  
XX Feravidobacterium pennavorans Ven 5 thermostable pullulanase DNA.  
XX  
XX Feravidobacterium pennavorans; Ven 5; thermostable pullulanase; pulla;  
XX Industrial saccharification; ss.  
XX  
XX Feravidobacterium pennavorans.  
XX  
XX WO9935274-A2.  
XX  
XX 15-JUL-1999.  
XX  
XX PD

XX  
XX 05-JAN-1999; 99WO-IB00069.  
XX  
XX 07-JAN-1998; 98US-0003834.  
XX  
XX (NOVO) NOVO-NORDISK AS.  
XX  
XX Dufferer F, Jorgensen PL;  
XX  
XX WPI; 1999-430401/36.  
XX  
XX P-PSDB; AAU24380.  
XX  
XX Nucleic acid encoding Feravidobacterium sp. Ven 5 thermostable  
XX pullulanase, useful for industrial saccharification processes  
XX  
XX Claim 1; Page 40-41; 52pp; English.  
XX  
XX The present sequence encodes Feravidobacterium pennavorans Ven 5  
XX thermostable pullulanase. Thermostable pullulanases are useful in  
XX industrial saccharification processes (see EP63909).  
XX  
XX Sequence 2601 BP; 881 A; 494 C; 596 G; 630 T; 0 other;  
XX  
XX Query Match 72.5%; Score 17.4; DB 20; Length 2601;  
XX Best Local Similarity 94.7%; Pred. No. 1.4e+02;  
XX Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
XX  
XX 3 TTTCTTGCCAGAGCGCC 21  
XX ||||||||||||||||  
XX Db 1924 TTTCTTGCCAGAGCGCC 1906  
XX  
XX RESULT 9  
XX AAU84218/c  
XX ID AAU84218 standard; DNA; 36135 BP.  
XX  
XX AAU84218;  
XX  
XX 07-NOV-2001 (first entry)  
XX  
XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:39030.  
XX  
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;  
XX cytostatic; gene therapy; vaccine; metastasis; ds.  
XX  
XX Homo sapiens.  
XX  
XX WO200157182-A2.  
XX  
XX 09-AUG-2001.  
XX  
XX 17-JAN-2001; 2001WO-US01354.  
XX  
XX 31-JAN-2000; 2000US-0179065.  
XX  
XX 04-FEB-2000; 2000US-0180628.  
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XX 24-FEB-2000; 2000US-0184664.  
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XX 02-MAR-2000; 2000US-0186350.  
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XX 16-MAR-2000; 2000US-0189874.  
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XX 18-APR-2000; 2000US-0198123.  
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XX 19-MAY-2000; 2000US-0205515.  
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XX 07-JUN-2000; 2000US-0209467.  
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XX 30-JUN-2000; 2000US-0215135.  
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XX 07-JUL-2000; 2000US-0216647.  
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XX 07-JUL-2000; 2000US-0216880.  
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XX 14-AUG-2000; 2000US-0224519.  
XX  
XX PD

PR 14-AUG-2000; 2000US-0225213.  
 PR 14-AUG-2000; 2000US-0225214.  
 PR 14-AUG-2000; 2000US-0225265.  
 PR 14-AUG-2000; 2000US-0225267.  
 PR 14-AUG-2000; 2000US-0225268.  
 PR 14-AUG-2000; 2000US-0225270.  
 PR 14-AUG-2000; 2000US-0225447.  
 PR 14-AUG-2000; 2000US-0225757.  
 PR 14-AUG-2000; 2000US-0225758.  
 PR 14-AUG-2000; 2000US-0225759.  
 PR 18-AUG-2000; 2000US-0226279.  
 PR 22-AUG-2000; 2000US-0226681.  
 PR 22-AUG-2000; 2000US-0226868.  
 PR 23-AUG-2000; 2000US-0227182.  
 PR 30-AUG-2000; 2000US-0228924.  
 PR 01-SEP-2000; 2000US-0228287.  
 PR 01-SEP-2000; 2000US-0229343.  
 PR 01-SEP-2000; 2000US-0229344.  
 PR 05-SEP-2000; 2000US-0229509.  
 PR 05-SEP-2000; 2000US-0229513.  
 PR 06-SEP-2000; 2000US-0230437.  
 PR 06-SEP-2000; 2000US-0230438.  
 PR 08-SEP-2000; 2000US-0231242.  
 PR 08-SEP-2000; 2000US-0231243.  
 PR 08-SEP-2000; 2000US-0231244.  
 PR 08-SEP-2000; 2000US-0231413.  
 PR 08-SEP-2000; 2000US-0231414.  
 PR 08-SEP-2000; 2000US-0232080.  
 PR 12-SEP-2000; 2000US-0231968.  
 PR 14-SEP-2000; 2000US-0232397.  
 PR 14-SEP-2000; 2000US-0232398.  
 PR 14-SEP-2000; 2000US-0232399.  
 PR 14-SEP-2000; 2000US-0232400.  
 PR 14-SEP-2000; 2000US-0232401.  
 PR 14-SEP-2000; 2000US-0233063.  
 PR 14-SEP-2000; 2000US-0233064.  
 PR 21-SEP-2000; 2000US-0233065.  
 PR 21-SEP-2000; 2000US-0234223.  
 PR 21-SEP-2000; 2000US-0234274.  
 PR 25-SEP-2000; 2000US-0234597.  
 PR 25-SEP-2000; 2000US-0234998.  
 PR 25-SEP-2000; 2000US-0234998.  
 PR 27-SEP-2000; 2000US-0235844.  
 PR 27-SEP-2000; 2000US-0235834.  
 PR 27-SEP-2000; 2000US-0235836.  
 PR 29-SEP-2000; 2000US-0236327.  
 PR 29-SEP-2000; 2000US-0236327.  
 PR 29-SEP-2000; 2000US-0236367.  
 PR 29-SEP-2000; 2000US-0236368.  
 PR 29-SEP-2000; 2000US-0236369.  
 PR 02-OCT-2000; 2000US-0236370.  
 PR 02-OCT-2000; 2000US-0236802.  
 PR 02-OCT-2000; 2000US-0237037.  
 PR 02-OCT-2000; 2000US-0237038.  
 PR 02-OCT-2000; 2000US-0237039.  
 PR 03-OCT-2000; 2000US-0237040.  
 PR 13-OCT-2000; 2000US-0239935.  
 PR 20-OCT-2000; 2000US-0239937.  
 PR 20-OCT-2000; 2000US-0240960.  
 PR 20-OCT-2000; 2000US-0241221.  
 PR 20-OCT-2000; 2000US-0241785.  
 PR 20-OCT-2000; 2000US-0241786.  
 PR 20-OCT-2000; 2000US-0241787.  
 PR 20-OCT-2000; 2000US-0241787.  
 PR 20-OCT-2000; 2000US-0241808.  
 PR 20-OCT-2000; 2000US-0241809.  
 PR 01-NOV-2000; 2000US-0241826.  
 PR 08-NOV-2000; 2000US-0246474.  
 PR 08-NOV-2000; 2000US-0246475.  
 PR 08-NOV-2000; 2000US-0246476.  
 PR 08-NOV-2000; 2000US-0246477.  
 PR 08-NOV-2000; 2000US-0246478.

PR 08-NOV-2000; 2000US-0246523.  
 PR 08-NOV-2000; 2000US-0246524.  
 PR 08-NOV-2000; 2000US-0246525.  
 PR 08-NOV-2000; 2000US-0246526.  
 PR 08-NOV-2000; 2000US-0246527.  
 PR 08-NOV-2000; 2000US-0246528.  
 PR 08-NOV-2000; 2000US-0246532.  
 PR 08-NOV-2000; 2000US-0246609.  
 PR 08-NOV-2000; 2000US-0246610.  
 PR 08-NOV-2000; 2000US-0246611.  
 PR 17-NOV-2000; 2000US-0246613.  
 PR 17-NOV-2000; 2000US-0249207.  
 PR 17-NOV-2000; 2000US-0249208.  
 PR 17-NOV-2000; 2000US-0249209.  
 PR 17-NOV-2000; 2000US-0249210.  
 PR 17-NOV-2000; 2000US-0249211.  
 PR 17-NOV-2000; 2000US-0249212.  
 PR 17-NOV-2000; 2000US-0249213.  
 PR 17-NOV-2000; 2000US-0249214.  
 PR 17-NOV-2000; 2000US-0249215.  
 PR 17-NOV-2000; 2000US-0249216.  
 PR 17-NOV-2000; 2000US-0249217.  
 PR 17-NOV-2000; 2000US-0249218.  
 PR 17-NOV-2000; 2000US-0249244.  
 PR 17-NOV-2000; 2000US-0249245.  
 PR 17-NOV-2000; 2000US-0249264.  
 PR 17-NOV-2000; 2000US-0249265.  
 PR 17-NOV-2000; 2000US-0249297.  
 PR 17-NOV-2000; 2000US-0249299.  
 PR 17-NOV-2000; 2000US-0249300.  
 PR 01-DEC-2000; 2000US-0250160.  
 PR 01-DEC-2000; 2000US-0250391.  
 PR 05-DEC-2000; 2000US-0251030.  
 PR 05-DEC-2000; 2000US-0251988.  
 PR 05-DEC-2000; 2000US-0256719.  
 PR 06-DEC-2000; 2000US-0251479.  
 PR 08-DEC-2000; 2000US-0251856.  
 PR 08-DEC-2000; 2000US-0251856.  
 PR 08-DEC-2000; 2000US-0251868.  
 PR 08-DEC-2000; 2000US-0251869.  
 PR 08-DEC-2000; 2000US-0251889.  
 PR 08-DEC-2000; 2000US-0251990.  
 PR 11-DEC-2000; 2000US-0254097.  
 PR 05-JAN-2001; 2001US-0259678.  
 (HUMA-) HUMAN GENOME SCI INC.  
 Rosen CA, Barash SC, Ruben SM;  
 WPL; 2001-483426/52.  
 Nucleic acids encoding human immune/hematopoietic antigen polypeptides,  
 useful for preventing, diagnosing and/or treating cancers and  
 metastasis -  
 Disclosure: SEQ ID NO 39030; 3071pp + Sequence Listing; English.  
 AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)  
 amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic  
 activity, and can be used in gene therapy and vaccine production. (I)  
 proteins and polynucleotides may be used in the prevention, diagnosis and  
 treatment of diseases associated with inappropriate (I) expression. For  
 example, they may be used to treat disorders associated with decreased  
 expression by rectifying mutations or deletions in a patient's genome  
 that affect the activity of (I) by expressing inactive proteins or to  
 supplement the patient's own production of (I). Additionally, (I)  
 polynucleotides may be used to produce the secreted (I), by inserting  
 the nucleic acids into a host cell and culturing the cell to express the  
 protein. (I) proteins and polynucleotides may be used to prevent,  
 diagnose and treat immune/hematopoietic-related diseases, especially  
 cancers and cancer metastases of hematopoietic-derived cells. AAK64703  
 to AAK67694 represent human immune/hematopoietic antigen genomic  
 sequences from the present invention. AAK54942 to AAK54950 and AAM82169  
 represent sequences used in the exemplification of the present invention.

XX Sequence 36135 BP; 10321 A; 8075 C; 8063 G; 9676 T; 0 other;  
 SQ Query Match 70.8%; Score 17; DB 22; Length 36135;  
 Best Local Similarity 100.0%; Pred. No. 3.5e+02;  
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTCTTGGCAAGAGC 18  
 ||||||||||||||||  
 DB 920 CTTCTTGGCAAGAGC 904

RESULT 10  
 ABV12459  
 ID ABV12459 standard; cDNA; 467 BP.  
 AC ABV12459;  
 XX 13-SEP-2002 (first entry)  
 DT  
 XX Human prostate expression marker CDNA 12450.  
 DE  
 XX Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
 KW pharmacogenomic marker; gene; ss.  
 KM  
 OS Homo sapiens.  
 XX  
 XX WO200160860-A2.  
 PN  
 XX 23-AUG-2001.  
 PD  
 XX 20-FEB-2001; 2001WO-US05171.  
 PF  
 XX 17-FEB-2000; 2000US-183319P.  
 PR 16-MAR-2000; 2000US-189862P.  
 PR 25-MAY-2000; 2000US-207454P.  
 PR 09-JUN-2000; 2000US-211314P.  
 PR 18-JUL-2000; 2000US-219007P.  
 PR 13-DEC-2000; 2000US-255281P.

(MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
 PA Schlegel R, Endege WO, Monahan JE;  
 PI WPI; 2001-662795/76.  
 PS  
 XX Novel isolated nucleic acid molecule associated with cancerous state of  
 PT prostate cells and correlating with presence of prostate cancer, useful  
 PT for detecting presence of prostate cancer, stage of prostate cancer -  
 XX  
 XX Claim 1; Page 2053; 11750pp; English.

The invention relates to an isolated nucleic acid molecule (I) comprising  
 a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
 specification or its complement. (I) is useful for:  
 (a) assessing whether a patient is afflicted with prostate cancer;  
 (b) monitoring the progression of prostate cancer in a patient;  
 (c) assessing the efficacy of a test compound to inhibit prostate  
 cancer in a patient;  
 (d) assessing the efficacy of a therapy for inhibiting prostate cancer  
 in a patient;  
 (e) selecting a composition for inhibiting prostate cancer in a patient;  
 (f) assessing the prostate cell carcinogenic potential of a compound;  
 (g) determining whether prostate cancer has metastasized in a patient;  
 (h) assessing the aggressiveness or indolence of prostate cancer in a  
 patient;  
 (I) is also useful as a pharmacodynamic or pharmacogenomic marker.

Sequence 467 BP; 71 A; 115 C; 132 G; 149 T; 0 other;

Query Match 70.0%; Score 16.8; DB 23; Length 467;  
 Best Local Similarity 90.0%; Pred. No. 2.1e+02;  
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GCTTCTTGGCAAGAGCGC 20  
 |||||||||||||  
 DB 330 GCTTCTTGGCAAGAGCGC 349

RESULT 11  
 ABV33599  
 ID ABV33599 standard; cDNA; 506 BP.  
 AC ABV33599;  
 XX 16-SEP-2002 (first entry)  
 DT  
 XX Human prostate expression marker CDNA 33590.  
 DE  
 XX Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
 KW pharmacogenomic marker; gene; ss.  
 KM  
 OS Homo sapiens.  
 XX  
 XX WO200160860-A2.  
 PN  
 XX 23-AUG-2001.  
 PD  
 XX 20-FEB-2001; 2001WO-US05171.  
 PF  
 XX 17-FEB-2000; 2000US-183319P.  
 PR 16-MAR-2000; 2000US-189862P.  
 PR 25-MAY-2000; 2000US-207454P.  
 PR 09-JUN-2000; 2000US-211314P.  
 PR 18-JUL-2000; 2000US-219007P.  
 PR 13-DEC-2000; 2000US-255281P.

(MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
 PA Schlegel R, Endege WO, Monahan JE;  
 PI WPI; 2001-662795/76.  
 PS  
 XX Novel isolated nucleic acid molecule associated with cancerous state of  
 PT prostate cells and correlating with presence of prostate cancer, useful  
 PT for detecting presence of prostate cancer, stage of prostate cancer -  
 XX  
 XX Claim 1; Page 7105; 11750pp; English.

The invention relates to an isolated nucleic acid molecule (I) comprising  
 a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
 specification or its complement. (I) is useful for:  
 (a) assessing whether a patient is afflicted with prostate cancer;  
 (b) monitoring the progression of prostate cancer in a patient;  
 (c) assessing the efficacy of a test compound to inhibit prostate  
 cancer in a patient;  
 (d) assessing the efficacy of a therapy for inhibiting prostate cancer  
 in a patient;  
 (e) selecting a composition for inhibiting prostate cancer in a patient;  
 (f) assessing the prostate cell carcinogenic potential of a compound;  
 (g) determining whether prostate cancer has metastasized in a patient;  
 (h) assessing the aggressiveness or indolence of prostate cancer in a  
 patient;  
 (I) is also useful as a pharmacodynamic or pharmacogenomic marker.

Sequence 506 BP; 80 A; 129 C; 145 G; 152 T; 0 other;

Query Match 70.0%; Score 16.8; DB 23; Length 506;  
 Best Local Similarity 90.0%; Pred. No. 2.1e+02;  
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GCTTCTTGGCAAGAGCGC 20  
 |||||||||||||  
 DB 376 GCTTCTTGGCAAGAGCGC 395

RESULT 12  
 ABK63732  
 ID ABK63732 standard; cDNA; 1679 BP.  
 XX  
 AC ABK63732;  
 XX  
 DT 18-JUN-2002 (first entry)  
 XX  
 DE Rat sequence differentially expressed in response to a hepatotoxin #1639.  
 XX  
 KW Rat; ss; hepatotoxin; expressed sequence tag; EST; drug screening;  
 KM differential expression; centrilobular necrosis; steatosis.  
 XX  
 OS Rattus norvegicus.  
 XX  
 PN W0200210453-A2.  
 XX  
 PD 07-FEB-2002.  
 XX  
 PE 30-JUL-2001; 2001WO-US23872.  
 XX  
 PR 31-JUL-2000; 2000US-222040P.  
 XX  
 PR 02-NOV-2000; 2000US-244880P.  
 PR 11-MAY-2001; 2001US-290022P.  
 PR 15-MAY-2001; 2001US-290645P.  
 PR 22-MAY-2001; 2001US-292336P.  
 PR 06-JUN-2001; 2001US-295798P.  
 PR 13-JUN-2001; 2001US-297457P.  
 PR 19-JUN-2001; 2001US-298884P.  
 PR 09-JUL-2001; 2001US-303455P.  
 XX  
 PA (GENE-) GENE LOGIC INC.  
 XX  
 PI Mendrick D, Porter MW, Johnson KR, Castle AL, Elashoff MR;  
 XX  
 DR WPI; 2002-241625/29.  
 XX  
 PT Predicting toxic effects of compounds or the progression of these toxic  
 PT effects by determining the changes in gene expression in tissues or  
 PT cells exposed to the toxin and comparing these to gene expression in  
 PT unexposed tissues or cells -  
 XX  
 PS Claim 1; Seq ID No 1639; 239pp; English.  
 XX  
 CC The invention relates to methods for predicting toxic effects of  
 CC compounds or the progression of these toxic effects by determining the  
 CC global changes in gene expression in tissues or cells exposed to the  
 CC toxin and comparing these to gene expression in unexposed tissues or  
 CC cells. Also included are methods of predicting at least one toxic  
 CC effect of a compound or progression of a toxic effect, preferably the  
 CC hepatotoxicity of a compound, comprising detecting the level of  
 CC expression in a tissue or cell sample exposed to the compound of two or  
 CC more genes listed in the specification, where differential expression of  
 CC the genes is indicative of at least one toxic effect or progression.  
 CC The method can also be used to identify an agent which modulates the  
 CC toxic response and predict cellular pathways that a compound modulates  
 CC in a cell. The methods utilize a set of at least two probes (on a solid  
 CC support in kit form), where each of the probes comprises a sequence that  
 CC specifically hybridizes to a gene listed in the specification, a computer  
 CC system comprising a database containing information identifying the  
 CC expression level in a tissue or cell sample exposed to a hepatotoxin of a  
 CC set of genes comprising at least two genes listed in the specification, a  
 CC and a user interface to view the information used to present information,  
 CC identifying the expression level in a tissue or cell of at least one gene  
 CC listed in the specification. The method is useful for elucidating global  
 CC changes in gene expression and for identifying toxicity markers in  
 CC tissues or cell exposed to a known toxin. The genes may be used as  
 CC toxicity markers in drug screening and toxicity assays. The genes and  
 CC gene expression information may be used as diagnostic markers for the  
 CC prediction or identification of the physiological state of tissue or cell  
 CC sample that has been exposed to a compound or agent. Hepatotoxicity  
 CC is characterized by centrilobular necrosis and steatosis. The present  
 CC sequence is an expressed sequence tag (EST) or cDNA derived from a gene

CC which is differentially expressed in response to a hepatotoxic agent.  
 XX  
 SO Sequence 1679 BP; 377 A; 471 C; 483 G; 348 T; 0 other;  
 XX  
 Query Match 70.0%; Score 16.8; DB 24; Length 1679;  
 Best Local Similarity 90.0%; Pred. NO. 2.6e+02;  
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 XX  
 QY 5 TCTTGGCCAGAGCGCCGCA 24  
 ||||||| |||||||  
 Db 61 TCTTGGCCAGAGCGCCGCA 80  
 XX  
 RESULT 13  
 ABL02991  
 ID ABL02991 standard; cDNA; 2304 BP.  
 XX  
 AC ABL02991;  
 XX  
 DT 26-MAR-2002 (first entry)  
 XX  
 DE Drosophila melanogaster expressed polynucleotide SEQ ID NO 3455.  
 XX  
 KW Drosophila; developmental biology; cell signalling; insecticide;  
 KM pharmaceutical; gene; ss.  
 XX  
 OS Drosophila melanogaster.  
 XX  
 PN W0200171042-A2.  
 XX  
 PD 27-SEP-2001.  
 XX  
 PE 23-MAR-2001; 2001WO-US09221.  
 XX  
 PR 23-MAR-2000; 2000US-191637P.  
 PR 11-JUL-2000; 2000US-0614150.  
 XX  
 PA (PEKE ) PE CORP NY.  
 XX  
 PI Venter JC, Adams M, Li PWD, Myers EW;  
 XX  
 DR WPI; 2001-656860/75.  
 DR P-PSDB; ABB58888.  
 XX  
 PT New isolated nucleic acid detection reagent for detecting 1000 or more  
 PT genes from Drosophila and for elucidating cell signalling and cell-cell  
 PT interactions -  
 XX  
 PS Claim 1; SEQ ID NO 3455; 21pp + Sequence Listing; English.  
 XX  
 CC The invention relates to an isolated nucleic acid detection reagent  
 CC capable of detecting 1000 or more genes from Drosophila. The invention is  
 CC useful in developmental biology and in elucidating cell signalling and  
 CC cell-cell interactions in higher eukaryotes for the development of  
 CC insecticides, therapeutics and pharmaceutical drugs. The invention  
 CC discloses genomic DNA sequences (AB101840-AB16175) and the encoded proteins  
 CC (AB57737-AB72072).  
 CC The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pcl\_sequences.  
 XX  
 SO Sequence 2304 BP; 541 A; 640 C; 721 G; 402 T; 0 other;  
 XX  
 Query Match 70.0%; Score 16.8; DB 23; Length 2304;  
 Best Local Similarity 90.0%; Pred. NO. 2.7e+02;  
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 XX  
 QY 4 TCTTTGCCAGAGCGCCGC 23  
 ||||||| |||||||  
 Db 226 TCTTTGCCAGAGCGCCGC 245

RESULT 14  
ABL02990  
ID ABL02990 standard; cDNA; 4446 BP.  
XX  
XX ABL02990;  
AC  
XX  
XX 26-MAR-2002 (first entry)  
DT  
XX  
XX Drosophila melanogaster expressed polynucleotide SEQ ID NO 3452.  
DE  
XX  
XX Drosophila; developmental biology; cell signalling; insecticide;  
KM  
XX  
XX pharmaceutical; gene; ss.  
OS  
XX  
XX Drosophila melanogaster.  
PN  
XX  
XX WO200171042-A2.  
PD  
XX  
XX 27-SEP-2001.  
PF  
XX  
XX 23-MAR-2001; 2001WO-US09231.  
PR  
XX  
XX 23-MAR-2000; 2000US-191637P.  
PS  
XX  
XX 11-JUL-2000; 2000US-0614150.  
XX  
XX (PEKE ) PE CORP NY.  
XX  
XX Venter JC, Adams M, Li PWD, Myers EW;  
PI  
XX  
XX WPI: 2001-656860/75.  
DR  
XX  
XX P-PSDB; ABB58887.  
XX  
XX  
XX New isolated nucleic acid detection reagent for detecting 1000 or more  
PT genes from Drosophila and for elucidating cell signalling and cell-cell  
PT interactions -  
PS  
XX  
XX Claim 1: SEQ ID NO 3452; 21pp + Sequence Listing; English.  
XX  
XX The invention relates to an isolated nucleic acid detection reagent  
XX capable of detecting 1000 or more genes from Drosophila. The invention is  
XX useful in developmental biology and in elucidating cell signalling and  
XX cell-cell interactions in higher eukaryotes for the development of  
XX insecticides, therapeutics and pharmaceutical drugs. The invention  
XX discloses genomic DNA sequences (AB116176-AB130511), expressed DNA  
XX sequences (AB101840-AB116175) and the encoded proteins  
XX (AB57737-AB572072).  
XX The sequence data for this patent did not form part of the printed  
XX specification, but was obtained in electronic format directly from WIPO  
XX at ftp.wipo.int/pub/published\_pcl\_sequences.  
SQ  
XX  
XX Sequence 4446 BP; 1191 A; 1023 C; 1084 G; 1148 T; 0 other;  
Query Match 70.0%; Score 16.8; DB 23; Length 4446;  
Best Local Similarity 90.0%; Pred. No. 3e+02; 2; Indels 0; Gaps 0;  
Matches 18; Conservative 0; Mismatches 0;

QY 4 TTCTTTCAGAGCGCGC 23  
DB 613 TTCTTTCAGAGCGCGC 632

RESULT 15  
ABK80729  
ID ABK80729 standard; DNA; 408 BP.  
XX  
XX ABK80729;  
AC  
XX  
XX 13-AUG-2002 (first entry)  
DT  
XX  
XX Bacillus clausii genomic sequence tag (GST) #3572.  
DE  
XX  
XX Differential gene expression; genomic sequenced tag; GST;  
KM altered culture condition; environmental stress;  
KW physiological provocation; ds.

XX  
XX Bacillus clausii.  
OS  
XX  
XX WO200229113-A2.  
PN  
XX  
XX 11-APR-2002.  
PD  
XX  
XX 05-OCT-2001; 2001WO-US31437.  
PF  
XX  
XX 06-OCT-2000; 2000US-0680598.  
PR  
XX  
XX 27-MAR-2001; 2001US-279526P.  
PS  
XX  
XX (NOVO ) NOVOZYMES BIOTECH INC.  
XX  
XX (NOVO ) NOVOZYMES AS.  
PA  
XX  
XX Berka R, Clausen IG;  
PI  
XX  
XX WPI: 2002-416684/44.  
DR  
XX  
XX  
XX Monitoring differential expression of several genes in first Bacillus  
PT cell relative to expression of same genes in one or more second  
PT Bacillus cells, by using substrate containing Bacillus genomic  
PT sequenced tag array -  
PS  
XX  
XX Claim 11: SEQ ID NO 8020; 200pp; English.  
XX  
XX The invention describes a method of monitoring differential expression of  
XX genes in a first Bacillus cell relative to expression of the genes in  
XX other Bacillus cells, comprising hybridising labelled nucleic acid probes  
XX isolated from Bacillus cells to a substrate containing array of Bacillus  
XX genomic sequenced tags (GST), examining the array, and determining  
XX relative gene expression by an observed hybridisation reporter signal of  
XX a spot in the array. The method is useful for measuring the expression of  
XX genes in a first Bacillus cell relative to expression of the same genes  
XX in one or more second Bacillus cells. The method is useful for monitoring  
XX global expression of several genes from a Bacillus cell, discovering new  
XX genes, identifying possible functions of unknown open reading frames and  
XX monitoring gene copy number variation and stability. Monitoring changes  
XX in expression of genes may be used to provide a representation of the way  
XX in which Bacillus cells adapt to changes in culture conditions.  
XX environmental stress or other physiological provocation. Extensive  
XX follow-up characterisation is unnecessary, when one spot on an array  
XX equals one gene or one open reading frame, since sequence information is  
XX available. This sequence represents a genomic sequence tag (GST) used in  
XX the method of the invention.  
XX Note: The sequence data for this patent did not form part of the printed  
XX specification, but was obtained in electronic format directly from WIPO  
XX at  
XX ftp.wipo.int/pub/published\_pcl\_sequences.  
SQ  
XX  
XX Sequence 408 BP; 106 A; 86 C; 104 G; 111 T; 1 other;  
Query Match 69.2%; Score 16.6; DB 24; Length 408;  
Best Local Similarity 82.6%; Pred. No. 2.5e+02; 4; Indels 0; Gaps 0;  
Matches 19; Conservative 0; Mismatches 4;

QY 1 GCTTTTTCAGAGCGCGC 23  
DB 374 GCATTCTTTCAGAGCGCGC 396

Search completed: March 17, 2003, 10:50:45  
Job time : 146.715 secs

GenCore version 5.1.4.p5\_4578  
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OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 10:23:12 ; Search time 848.387 Seconds

(without alignments)  
458.154 Million cell updates/sec

Title: US-09-836-439-3

Perfect score: 24  
Sequence: 1 gcttccttgccagagcgccgca 24

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Maximum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database :

EST:  
1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estnu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hic:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hic:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: gb\_gss:\*  
18: em\_gss\_hum:\*  
19: em\_gss\_inv:\*  
20: em\_gss\_pln:\*  
21: em\_gss\_vrt:\*  
22: em\_gss\_fun:\*  
23: em\_gss\_mam:\*  
24: em\_gss\_mus:\*  
25: em\_gss\_other:\*  
26: em\_gss\_pro:\*  
27: em\_gss\_rtd:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	21.4	89.2	303	14	BM694379
2	21.4	89.2	319	14	BM637565
3	21.4	89.2	337	14	BM723222
4	21.4	89.2	340	14	BM682444
5	21.4	89.2	349	14	BM703950
6	21.4	89.2	441	14	BM690151

Result 1	LOCUS	DEFINITION	ACCESSION	VERSION	KEYWORDS	SOURCE	ORGANISM	REFERENCE	AUTHORS	TITLE	JOURNAL	MEDLINE	COMMENT
BM694379	303 bp	UI-E-C11-afp-p-18-0-UI-r1 UI-E-C11 Homo sapiens cDNA clone	BM694379	1	GI:19007637	human.	Homo sapiens	1	Bonaldi, M.F., Lennon, G. and Soares, M.B.	Normalization and subtraction: two approaches to facilitate gene discovery	Genome Res. 6 (9), 791-806 (1996)	97044477	Contact: Soares, MB Program for Rat Gene Discovery and Mapping University of Iowa 451 Eckstein Medical Research Building Iowa City, IA 52242, USA Tel: 319 335 8250 Fax: 319 335 9565 Email: msoares@blue.weeg.uiowa.edu Tissue Procurement: Dr. Gregg Hageman CDNA library preparation: Dr. M. Bento Soares, University of Iowa CDNA library arrayed by: Dr. M. Bento Soares, University of Iowa DNA Sequencing by: Dr. W. Bento Soares, University of Iowa

## ALIGNMENTS

7	21.4	89.2	442	9	AL712412	AL712412 DKFZ6860
8	21.4	89.2	446	14	BM696193	BM696193 UI-E-C11-
9	21.4	89.2	469	14	BM688227	BM688227 UI-E-C10-
10	21.4	89.2	471	9	AL712402	AL712402 DKFZ686N
11	21.4	89.2	471	14	B0250368	B0250368 TAE25006A
12	21.4	89.2	474	14	BM688059	BM688059 UI-E-C10-
13	21.4	89.2	478	13	BM662914	BM662914 UI-E-C10-
14	21.4	89.2	493	13	BM690137	BM690137 UI-E-C10-
15	21.4	89.2	493	14	BM691592	BM691592 UI-E-C11-
16	21.4	89.2	496	14	BM690311	BM690311 UI-E-C10-
17	21.4	89.2	499	14	B0640665	B0640665 hb31f09.y
18	21.4	89.2	503	14	B0636862	B0636862 hb02e01.y
19	21.4	89.2	504	14	BM686188	BM686188 UI-E-C10-
20	21.4	89.2	505	14	T27877	T27877 EST19137 Hu
21	21.4	89.2	510	14	B0637734	B0637734 hb13h04.y
22	21.4	89.2	513	14	BM690319	BM690319 UI-E-C10-
23	21.4	89.2	514	14	B0640660	B0640660 hb31f02.y
24	21.4	89.2	526	14	B0639829	B0639829 hb20h01.y
25	21.4	89.2	530	14	BM690115	BM690115 UI-E-C10-
26	21.4	89.2	530	14	B0638124	B0638124 hb18g07.y
27	21.4	89.2	535	14	B0638268	B0638268 hb20e05.y
28	21.4	89.2	535	14	B0640241	B0640241 hb26e01.y
29	21.4	89.2	549	14	B0636918	B0636918 hb03c02.y
30	21.4	89.2	549	14	B0639356	B0639356 hb14e06.y
31	21.4	89.2	562	14	B0636602	B0636602 hb11h09.y
32	21.4	89.2	575	14	B0640490	B0640490 hb29d06.y
33	21.4	89.2	576	14	BM704430	BM704430 UI-E-C11-
34	21.4	89.2	578	14	BM685757	BM685757 UI-E-C10-
35	21.4	89.2	578	14	BM688176	BM688176 UI-E-C10-
36	21.4	89.2	580	14	BM688912	BM688912 UI-E-C10-
37	21.4	89.2	587	14	B0637208	B0637208 hb07a10.y
38	21.4	89.2	590	14	B0640394	B0640394 hb27h12.y
39	21.4	89.2	591	9	AL711507	AL711507 DKFZ686D
40	21.4	89.2	595	14	B0638291	B0638291 hb20e07.y
41	21.4	89.2	599	14	B0638906	B0638906 hb03a11.y
42	21.4	89.2	604	14	B0638120	B0638120 hb18g02.y
43	21.4	89.2	605	9	AL712251	AL712251 DKFZ686L
44	21.4	89.2	606	14	B0638554	B0638554 hb24h07.y
45	21.4	89.2	606	14	B0639637	B0639637 hb18a09.y

Clone Distribution: Researchers may obtain clones from Research Genetics (www.resgen.com).

Seq primer: M13 Reverse.

#### FEATURES

##### Source

Location/Qualifiers

1. .303

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="UI-E-C11-afp-p-18-0-UI"

/clone\_lib="UI-E-C11"

/tissue\_type="RPE and Choroid"

/dev\_stage="adult"

/lab\_host="DH10B (Life Technologies) (T1 phage resistant)"

/note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a modified polylinker; Site: 1; Ecor I; Site: 2; Not I; UI-E-C11 is a normalized cDNA library containing the following tissue(s): RPE and Choroid. The library was constructed according to Bonaldi, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an Ecor I adaptor, digested with Not I, and cloned directionally into pT73-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is ACCCTA. This library was created for the program, Gene Discovery in the Visual system, supported by National Eye Institute (NEI)."

#### BASE COUNT

70 a 101 c 67 g 65 t

#### ORIGIN

Query Match

Best Local Similarity 95.2%; Score 21.4; DB 14; Length 303;

Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTGGCAAGAGCGCGC 23

Db 171 GCGTCTTGGCAAGAGCGCGC 193

#### RESULT 2

##### LOCUS

B0637565 319 bp mRNA linear EST 15-JUL-2002

Definition hel1904.y1 Human Retina cDNA (Un-normalized, unamplified): hd/he

##### ACCESSION

B0637565

##### VERSION

##### KEYWORDS

##### SOURCE

##### ORGANISM

##### REFERENCE

##### AUTHORS

##### TITLE

##### JOURNAL

##### COMMENT

##### FEATURES

##### Source

##### Location/Qualifiers

##### 1. .319

##### /organism="Homo sapiens"

##### /db\_xref="taxon:9606"

##### /clone="hel1904"

/clone\_lib="Human Retina cDNA (Un-normalized, unamplified)

: hd/he"

/tissue\_type="Retina"

/dev\_stage="Adult"

/lab\_host="EMDH10B"

/note="Organ: Eye; Vector: pSPORT1; Neural retina tissue was dissected from two 80 year old donors with no observed eye disease. 100ug of total RNA was used for library construction. A directionally cloned cDNA library in the pSPORT1 vector (Life Technologies) was constructed at Bioserve Biotechnology (Laurel MD) essentially following the protocols of the SuperScript Plasmid System full details of which are contained in the manufacturer's instruction manual (http://www.lifetech.com/). First strand synthesis was carried out using a Not I primer-adaptor [5'-pGACATGTTCTAGATGCGAGCGCGC(7)15-3']

]. EST analysis was performed on the unamplified library at the NIH Intramural Sequencing Center (NISC)."

#### BASE COUNT

72 a 108 c 75 g 64 t

#### ORIGIN

Query Match

Best Local Similarity 95.7%; Score 21.4; DB 14; Length 319;

Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTGGCAAGAGCGCGC 23

Db 204 GCGTCTTGGCAAGAGCGCGC 226

#### RESULT 3

##### LOCUS

BM723222 337 bp mRNA linear EST 01-MAR-2002

Definition UI-E-EJ0-alo-1-15-0-UI.r1 UI-E-EJ0 Homo sapiens cDNA clone

##### ACCESSION

##### VERSION

##### KEYWORDS

##### SOURCE

##### ORGANISM

##### REFERENCE

##### AUTHORS

##### TITLE

##### JOURNAL

##### MEDLINE

##### COMMENT

##### FEATURES

##### Source

##### Location/Qualifiers

##### 1. .337

##### /organism="Homo sapiens"

##### /db\_xref="taxon:9606"

##### /clone="UI-E-EJ0-alo-1-15-0-UI"

##### /clone\_lib="UI-E-EJ0"

##### /tissue\_type="fetal eyes, lens, eye anterior segment, optic nerve, retina, Retina foveal and Macular, RPE and Choroid"

##### /dev\_stage="adult"

##### /lab\_host="EMDH10B"

##### /note="Organ: Eye; Vector: pSPORT1; Neural retina tissue was dissected from two 80 year old donors with no observed eye disease. 100ug of total RNA was used for library construction. A directionally cloned cDNA library in the pSPORT1 vector (Life Technologies) was constructed at Bioserve Biotechnology (Laurel MD) essentially following the protocols of the SuperScript Plasmid System full details of which are contained in the manufacturer's instruction manual (http://www.lifetech.com/). First strand synthesis was carried out using a Not I primer-adaptor [5'-pGACATGTTCTAGATGCGAGCGCGC(7)15-3']

##### ]. EST analysis was performed on the unamplified library at the NIH Intramural Sequencing Center (NISC)."

##### at the NIH Intramural Sequencing Center (NISC)."

##### at the NIH Intramural Sequencing Center (NISC)."

##### at the NIH Intramural Sequencing Center (NISC)."



```

/dev.stage="fetal and adult"
/lab.host="DH10B (Life Technologies) (T1 phage resistant)"
/notes="Organ: eye; Vector: pT73-Pac (Pharmacia) with a
modified polylinker; Site-1: EcoR I; Site-2: Not I;
UI-E-EJ0 is a subtracted cDNA library constructed
according to Bonaldo, Lennon and Soares, Genome Research,
6:791-806, 1996. First strand cDNA synthesis was primed
with an oligo-dT primer containing a Not I site. Double
stranded cDNA was ligated to an EcoR I adaptor, digested
with Not I, and cloned directionally into pT73-Pac
vector. The oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tags for this library are: fetal eyes, AGAATCAACA
; lens, CGATTAGCGA; eye anterior segment, AATGCCGCT;
optic nerve, CCAATACAG; retina, CCGCG; Retina foveal and
Macular, GTCC; RPE and Choroid, ACCTA. This library was
created for the program, Gene Discovery in the Visual
System, supported by National Eye Institute (NEI)."
```

BASE COUNT 86 a 105 c 74 g 72 t

ORIGIN

Query Match 89.2%; Score 21.4; DB 14; Length 337;  
Best Local Similarity 95.7%; Pred. No. 24;  
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTTCGCAAGAGCGCCG 23  
|||  
|||

Db 1 GCGTCTTTCGCAAGAGCGCCG 23

RESULT 4  
LOCUS BM682444 340 bp mRNA linear EST 27-FEB-2002  
DEFINITION UI-E-EJ0-a10-1-15-0-UI-s1 UI-E-EJ0 Homo sapiens cDNA clone  
ACCESSION BM682444  
VERSION BM682444.1 GI:18992340  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 340)  
Bonaldo,M.F., Lennon,G. and Soares,M.B.  
Normalization and subtraction: two approaches to facilitate gene  
discovery  
Genome Res. 6 (9), 791-806 (1996)  
97044477  
Contact: Soares, MB  
Program for Rat Gene Discovery and Mapping  
University of Iowa  
451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: mssoares@blue.weeg.uiowa.edu  
Tissue Procurement: Dr. Gregg Hageman  
cDNA Library preparation: Dr. M. Bento Soares, University of Iowa  
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
Clone Distribution: Researchers may obtain clones from Research  
Genetics (www.resgen.com).  
The following repetitive elements were found in this cDNA  
sequence: 1-37, >AT-rich#low\_complexity (matched complement)  
Seq primer: M13 Forward  
POLY-A=yes.

FEATURES  
source  
1..340  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="UI-E-EJ0-a10-1-15-0-UI"  
/clone\_lib="UI-E-EJ0"  
/tissue\_type="fetal eyes, lens, eye anterior segment,

```

optic nerve, retina, Retina foveal and Macular, RPE and
Choroid"
/dev.stage="fetal and adult"
/lab.host="DH10B (Life Technologies) (T1 phage resistant)"
/notes="Organ: eye; Vector: pT73-Pac (Pharmacia) with a
modified polylinker; Site-1: EcoR I; Site-2: Not I;
UI-E-EJ0 is a subtracted cDNA library constructed
according to Bonaldo, Lennon and Soares, Genome Research,
6:791-806, 1996. First strand cDNA synthesis was primed
with an oligo-dT primer containing a Not I site. Double
stranded cDNA was ligated to an EcoR I adaptor, digested
with Not I, and cloned directionally into pT73-Pac
vector. The oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tags for this library are: fetal eyes, AGAATCAACA
; lens, CGATTAGCGA; eye anterior segment, AATGCCGCT;
optic nerve, CCAATACAG; retina, CCGCG; Retina foveal and
Macular, GTCC; RPE and Choroid, ACCTA. This library was
created for the program, Gene Discovery in the Visual
System, supported by National Eye Institute (NEI)."
```

BASE COUNT 71 a 76 c 105 g 88 t

ORIGIN

Query Match 89.2%; Score 21.4; DB 14; Length 340;  
Best Local Similarity 95.7%; Pred. No. 24;  
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTTCGCAAGAGCGCCG 23  
|||  
|||

Db 331 GCGTCTTTCGCAAGAGCGCCG 309

RESULT 5  
LOCUS BM703950 349 bp mRNA linear EST 28-FEB-2002  
DEFINITION UI-E-CK1-afk-m-09-0-UI-r1 UI-E-CK1 Homo sapiens cDNA clone  
ACCESSION BM703950  
VERSION BM703950.1 GI:19017208  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 349)  
Bonaldo,M.F., Lennon,G. and Soares,M.B.  
Normalization and subtraction: two approaches to facilitate gene  
discovery  
Genome Res. 6 (9), 791-806 (1996)  
97044477  
Contact: Soares, MB  
Program for Rat Gene Discovery and Mapping  
University of Iowa  
451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: mssoares@blue.weeg.uiowa.edu  
Tissue Procurement: Dr. Gregg Hageman  
cDNA Library preparation: Dr. M. Bento Soares, University of Iowa  
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
Clone Distribution: Researchers may obtain clones from Research  
Genetics (www.resgen.com).  
Seq primer: M13 Reverse.

FEATURES  
source  
1..349  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="UI-E-CK1-afk-m-09-0-UI"

```

/clone_lib="UI-E-CK1"
/tissue_type="Retina Foveal and Macular"
/dev_stage="adult"
/lab_host="DH10B (Life Technologies) (T1 phage resistant)"
/Note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a
modified polylinker; Site_1: EcoR I; Site_2: Not I;
UI-E-CK1 is a normalized cDNA library containing the
following tissue(s): Retina Foveal and Macular. The
library was constructed according to Bonaldo, Lennon and
Soares, Genome Research, 6:791-806, 1996. First strand
cDNA synthesis was primed with an oligo-dT primer
containing a Not I site. Double stranded cDNA was ligated
to an EcoR I adaptor, digested with Not I, and cloned
directionally into pT73-Pac vector. The oligonucleotide
used to prime the synthesis of first-strand cDNA contains
a library tag sequence that is located between the Not I
site and the (dT)18 tail. The sequence tag for this
library is GTCC. This library was created for the program,
Gene Discovery in the Visual System, supported by National
Eye Institute (NEI)."
BASE COUNT      96 a      116 c      61 t
ORIGIN
Query Match      89.2%; Score 21.4; DB 14; Length 349;
Best Local Similarity 95.7%; Pred. No. 24;
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 GCTTCTTGGCCAGAGCGCCGC 23
    |||
Db 37 GCGTCTTGGCCAGAGCGCCGC 59

RESULT 6
LOCUS      BM690151      441 bp      mRNA      linear      EST 28-FEB-2002
DEFINITION UI-E-CK10-acd-f-11-0-UI.r1 UI-E-CK10 Homo sapiens cDNA clone
VERSION     BM690151
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE   1 (bases 1 to 441)
AUTHORS    Bonaldo,M.F., Lennon,G. and Soares,M.B.
TITLE      Normalization and subtraction: two approaches to facilitate gene
JOURNAL    Genome Res. 6 (9), 791-806 (1996)
MEDLINE    97044477
COMMENT     Contact: Soares, MB
            Program for Rat Gene Discovery and Mapping
            University of Iowa
            451 Eckstein Medical Research Building Iowa City, IA 52242, USA
            Tel: 319 335 8250
            Fax: 319 335 9565
            Email: msoares@blue.weeg.uiowa.edu
            Tissue Procurement: Dr. Gregg Hageman
            cDNA Library Preparation: Dr. K. Bento Soares, University of Iowa
            cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
            DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
            Clone Distribution: Researchers may obtain clones from Research
            Genetics (www.resgen.com).
            Seq primer: M13 Reverse
            Location/Qualifiers
                1..441
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="UI-E-CK10-acd-f-11-0-UI"
                /clone_lib="UI-E-CK10"
                /tissue_type="human retina"
                /dev_stage="adult"
                /lab_host="DH10B (Life Technologies) (T1 phage resistant)"

```

```

/Note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a
modified polylinker; Site_1: EcoR I; Site_2: Not I;
UI-E-CK10 is a cDNA library containing the following
tissue(s): retina. The library was constructed according
to Bonaldo, Lennon and Soares, Genome Research, 6:791-806,
1996. First strand cDNA synthesis was primed with an
oligo-dT primer containing a Not I site. Double stranded
cDNA was ligated to an EcoR I adaptor, digested with Not
I, and cloned directionally into pT73-Pac vector. The
oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tag for this library is CCGCG. This library was
created for the program, Gene Discovery in the Visual
System, supported by National Eye Institute (NEI)."
BASE COUNT      97 a      143 c      100 g      101 t
ORIGIN
Query Match      89.2%; Score 21.4; DB 14; Length 441;
Best Local Similarity 95.7%; Pred. No. 26;
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

```

```

Oy 1 GCTTCTTGGCCAGAGCGCCGC 23
    |||
Db 301 GCGTCTTGGCCAGAGCGCCGC 323

```

```

RESULT 7
LOCUS      AL712412      442 bp      mRNA      linear      EST 22-MAR-2002
DEFINITION DKFZp686O1888.r1 666 (synonym: nlcc3) Homo sapiens cDNA clone
VERSION     DKFZp686O1888.5', mRNA sequence.
AL712412
AL712412.1 GI:19695767
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE   1 (bases 1 to 442)
AUTHORS    Wambutt,R., Heubner,D., Mewes,W., Well,B. and Wiemann,S.
TITLE      EST (Wambutt,R., Heubner,D., Mewes,H.W., Well,B. and Wiemann,S.)
JOURNAL    Unpublished (1999)
COMMENT     Contact: Wambutt R
            MIPS
            Am Klopferspitz 18a D-82152 Martinsried, Germany
            This is the 5' sequence of the clone insert
            Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
            Research Center (DKFZ), Email s.wiemann@dkfz-heidelberg.de;
            sequenced by AGOWA (Berlin/Germany) within the cDNA sequencing
            consortium of the German Genome Project.
            No sl sequence available.
            This clone (DKFZp686O1888) is available at the RZPD in Berlin.
            Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
            Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.
            Location/Qualifiers
                1..442
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="DKFZp686O1888"
                /clone_lib="666 (synonym: nlcc3)"
                /tissue_type="human skeletal muscle"
                /dev_stage="adult"
                /lab_host="DH10B"
                /Note="Vector: pT73-Pac; Site_1: SfiIA; Site_2: SfiIB;
                cDNA collection"

```

```

BASE COUNT      106 a      144 c      97 g      95 t
ORIGIN
Query Match      89.2%; Score 21.4; DB 9; Length 442;
Best Local Similarity 95.7%; Pred. No. 26;
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

```

OY 1 GCTTCTTTGCCAAGAGCGCCGC 23  
 Db 23 GCCTTCTTTGCCAAGAGCGCCGC 45

# RESULT 8 BM696193

LOCUS 446 bp mRNA linear EST 28-FEB-2002

DEFINITION UI-E-CL1-afa-d-24-0-UI r1 UI-E-CL1 Homo sapiens cDNA clone

ACCESSION BM696193.1 GI:19009451

VERSION 1

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

Human.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 446)  
 Bonaldo,M.F., Lennon,G. and Soares,M.B.  
 Normalization and subtraction: two approaches to facilitate gene  
 discovery  
 Genome Res. 6 (9), 791-806 (1996)  
 Contact: Soares, MB  
 Program for Rat Gene Discovery and Mapping  
 University of Iowa  
 451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
 Tel: 319 335 8250  
 Fax: 319 335 9565  
 Email: msoares@blue.weeg.uiowa.edu  
 Tissue Procurement: Dr. Gregg Hageman  
 CDNA Library Preparation: Dr. M. Bento Soares, University of Iowa  
 DNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa  
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
 Clone Distribution: Researchers may obtain clones from Research  
 Genetics (www.resgen.com).  
 Seq primer: M13 Reverse.  
 Location/Qualifiers

## FEATURES

Source

1. 446  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone="UI-E-CL1-afa-d-24-0-UI"  
 /clone\_lib="UI-E-CL1"  
 /tissue\_type="human retina"  
 /dev\_stage="adult"  
 /lab\_host="PH10B (Life Technologies) (T1 phage resistant)"  
 /note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a  
 modified polylinker; Site\_1: EcoR I; Site\_2: Not I;  
 UI-E-CL1 is a normalized cDNA library containing the  
 following tissue(s): retina. The library was constructed  
 according to Bonaldo, Lennon and Soares, Genome Research,  
 6:791-806, 1996. First strand cDNA synthesis was primed  
 with an oligo-dT primer containing a Not I site. Double  
 stranded cDNA was ligated to an EcoR I adaptor, digested  
 with Not I, and cloned directionally into pT73-Pac  
 vector. The oligonucleotide used to prime the synthesis of  
 first-strand cDNA contains a library tag sequence that is  
 located between the Not I site and the (dT)18 tail. The  
 sequence tag for this library is CCGCG. This library was  
 created for the program, Gene Discovery in the Visual  
 System, supported by National Eye Institute (NEI)."  
 BASE COUNT 95 a 148 c 106 g 96 t 1 others

## Query Match

Best Local Similarity 89.2%; Score 21.4; DB 14; Length 446;  
 Pred. No. 26;

Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTTGCCAAGAGCGCCGC 23  
 Db 226 GCCTTCTTTGCCAAGAGCGCCGC 248

# RESULT 9 BM688227

LOCUS 469 bp mRNA linear EST 28-FEB-2002

DEFINITION UI-E-CL0-aby-g-03-0-UI r1 UI-E-CL0 Homo sapiens cDNA clone

ACCESSION BM688227.1 GI:19001478

VERSION 1

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

Human.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 469)  
 Bonaldo,M.F., Lennon,G. and Soares,M.B.  
 Normalization and subtraction: two approaches to facilitate gene  
 discovery  
 Genome Res. 6 (9), 791-806 (1996)  
 Contact: Soares, MB  
 Program for Rat Gene Discovery and Mapping  
 University of Iowa  
 451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
 Tel: 319 335 8250  
 Fax: 319 335 9565  
 Email: msoares@blue.weeg.uiowa.edu  
 Tissue Procurement: Dr. Gregg Hageman  
 CDNA Library Preparation: Dr. M. Bento Soares, University of Iowa  
 DNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa  
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
 Clone Distribution: Researchers may obtain clones from Research  
 Genetics (www.resgen.com).  
 The following repetitive elements were found in this cDNA  
 Sequence: 425-459, >AT-rich#low\_complexity  
 Seq primer: M13 Reverse.  
 Location/Qualifiers

## FEATURES

Source

1. 469  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone="UI-E-CL0-aby-g-03-0-UI"  
 /clone\_lib="UI-E-CL0"  
 /tissue\_type="human retina"  
 /dev\_stage="adult"  
 /lab\_host="PH10B (Life Technologies) (T1 phage resistant)"  
 /note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a  
 modified polylinker; Site\_1: EcoR I; Site\_2: Not I;  
 UI-E-CL0 is a cDNA library containing the following  
 tissue(s): retina. The library was constructed according  
 to Bonaldo, Lennon and Soares, Genome Research, 6:791-806,  
 1996. First strand cDNA synthesis was primed with an  
 oligo-dT primer containing a Not I site. Double stranded  
 cDNA was ligated to an EcoR I adaptor, digested with Not  
 I, and cloned directionally into pT73-Pac vector. The  
 oligonucleotide used to prime the synthesis of  
 first-strand cDNA contains a library tag sequence that is  
 located between the Not I site and the (dT)18 tail. The  
 sequence tag for this library is CCGCG. This library was  
 created for the program, Gene Discovery in the Visual  
 System, supported by National Eye Institute (NEI)."  
 BASE COUNT 110 a 155 c 99 g 104 t 1 others

## Query Match

Best Local Similarity 89.2%; Score 21.4; DB 14; Length 469;  
 Pred. No. 27;

Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTTGCCAAGAGCGCCGC 23  
 Db 82 GCCTTCTTTGCCAAGAGCGCCGC 104

## RESULT 10 AL712402

LOCUS 471 bp mRNA linear EST 22-MAR-2002



UI-E-ClO is a cDNA library containing the following tissue(s): retina. The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pRT3-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is CCGCG. This library was created for the program, Gene Discovery in the Visual System, supported by National Eye Institute (NEI).

BASE COUNT 110 a 158 c 105 g 101 t  
ORIGIN

Query Match 89.2%; Score 21.4; DB 14; Length 474;  
Best Local Similarity 95.7%; Pred. No. 27;  
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTTGCCAGAGCGCCG 23  
|||  
Db 41 GCGTCTTTGCCAGAGCGCCG 63

RESULT 13  
BM652914/c 478 bp mRNA linear EST 27-FEB-2002  
LOCUS UI-E-CKO-aan-e-04-0-UI.s1 UI-E-CKO Homo sapiens cDNA clone  
DEFINITION UI-E-CKO-aan-e-04-0-UI 3', mRNA sequence.  
ACCESSION BM652914  
VERSION BM652914.1 GI:18967816  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
AUTHORS 1 (bases 1 to 478)  
TITLE Normalization and subtraction: two approaches to facilitate gene discovery  
JOURNAL Genome Res. 6 (9), 791-806 (1996)  
MEDLINE 97044477  
COMMENT Contact: Soares, MB  
Program for Rat Gene Discovery and Mapping  
University of Iowa  
451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: msoares@blue.weeg.uiowa.edu  
Tissue Procurement: Dr. Gregg Hageman  
cDNA Library Preparation: Dr. M. Bento Soares, University of Iowa  
cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa  
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
Clone Distribution: Researchers may obtain clones from Research Genetics (www.resgen.com).  
Seq primer: M13 Forward  
POLYA=Yes.

#### FEATURES

Location/Qualifiers  
1..478  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="UI-E-CKO-aan-e-04-0-UI"  
/clone\_lib="UI-E-CKO"  
/tissue\_type="Retina Foveal and Macular"  
/dev\_stage="adult"  
/lab\_host="DH10B (Life Technologies) (T1 phage resistant)"  
/note="Organ: eye; Vector: pRT3-Pac (Pharmacia) with a modified polylinker; Site: 1: EcoR I; Site 2: Not I; UI-E-CKO is a cDNA library containing the following tissue(s): Retina Foveal and Macular. The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA

synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pRT3-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is CCGCG. This library was created for the program, Gene Discovery in the Visual System, supported by National Eye Institute (NEI).

TAG\_LIB-UI-E-CKO  
TAG\_TISSUE=Foveal and Macular Retina  
TAG\_SEQ=GTCC

BASE COUNT 86 a 109 c 157 g 123 t 3 others  
ORIGIN

Query Match 89.2%; Score 21.4; DB 13; Length 478;  
Best Local Similarity 95.7%; Pred. No. 27;  
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTTGCCAGAGCGCCG 23  
|||  
Db 307 GCGTCTTTGCCAGAGCGCCG 285

RESULT 14  
BM690137 493 bp mRNA linear EST 28-FEB-2002  
LOCUS UI-E-CKO-acd-d-08-0-UI.r1 UI-E-CKO Homo sapiens cDNA clone  
DEFINITION UI-E-CKO-acd-d-08-0-UI 5', mRNA sequence.  
ACCESSION BM690137  
VERSION BM690137.1 GI:19003395  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
AUTHORS 1 (bases 1 to 493)  
TITLE Normalization and subtraction: two approaches to facilitate gene discovery  
JOURNAL Genome Res. 6 (9), 791-806 (1996)  
MEDLINE 97044477  
COMMENT Contact: Soares, MB  
Program for Rat Gene Discovery and Mapping  
University of Iowa  
451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: msoares@blue.weeg.uiowa.edu  
Tissue Procurement: Dr. Gregg Hageman  
cDNA Library Preparation: Dr. M. Bento Soares, University of Iowa  
cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa  
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
Clone Distribution: Researchers may obtain clones from Research Genetics (www.resgen.com).  
The following repetitive elements were found in this cDNA sequence: 425-459, >AT-rich#low-complexity  
Seq primer: M13 Reverse.

#### FEATURES

Location/Qualifiers  
1..493  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="UI-E-CKO-acd-d-08-0-UI"  
/clone\_lib="UI-E-CKO"  
/tissue\_type="human retina"  
/dev\_stage="adult"  
/lab\_host="DH10B (Life Technologies) (T1 phage resistant)"  
/note="Organ: eye; Vector: pRT3-Pac (Pharmacia) with a modified polylinker; Site: 1: EcoR I; Site 2: Not I; UI-E-CKO is a cDNA library containing the following tissue(s): retina. The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806,

1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into p7773-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)<sub>18</sub> tail. The sequence tag for this library is CCGCG. This library was created for the program, Gene Discovery in the Visual System, supported by National Eye Institute (NEI)."

BASE COUNT 110 a 162 c 108 g 113 t  
ORIGIN

Query Match 89.2%; Score 21.4; DB 14; Length 493;  
Best Local Similarity 95.7%; Pred. No. 27;  
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
QY 1 GCTTCTTTGCCAAGAGCGCGC 23  
DB 131 GCGTCTTTGCCAAGAGCGCGC 153

RESULT 15 493 bp mRNA linear EST 28-FEB-2002  
BM691592  
LOCUS UI-E-C11-abh-c-07-0-UI.r1 UI-E-C11 Homo sapiens cDNA clone  
DEFINITION UI-E-C11-abh-c-07-0-UI 5', mRNA sequence.  
ACCESSION BM691592  
VERSION BM691592.1 GI:19004850  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.  
REFERENCE 1 (bases 1 to 493)  
AUTHORS Bonaldo,M.F., Lennon,G. and Soares,M.B.  
TITLE Normalization and subtraction: two approaches to facilitate gene  
discovery  
JOURNAL Genome Res. 6 (9), 791-806 (1996)  
MEDLINE 97044477  
COMMENT Contact: Soares, MB  
Program for Rat Gene Discovery and Mapping  
University of Iowa  
451 Eckstein Medical Research Building Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: msoares@blue.weeg.uiowa.edu  
Tissue Procurement: Dr. Gregg Hageman  
cDNA library preparation: Dr. M. Bento Soares, University of Iowa  
CDNA library Arrayed by: Dr. M. Bento Soares, University of Iowa  
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
Clone Distribution: Researchers may obtain clones from Research  
Genetics (www.resgen.com).  
The following repetitive elements were found in this cDNA  
sequence: 302-335, >AT-rich#Low\_complexity  
Seq primer: M13 Reverse.

# FEATURES

Location/Qualifiers  
1..493  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="UI-E-C11-abh-c-07-0-UI"  
/clone\_lib="UI-E-C11"  
/tissue\_type="RPE and Choroid"  
/dev\_stage="adult"  
/lab\_host="DH10B (Life Technologies) (T1 phage resistant)"  
/note="Organ: eye; Vector: p7773-Pac (Pharmacia) with a  
modified polylinker; Site\_1: EcoR I; Site\_2: Not I;  
UI-E-C11 is a normalized cDNA library containing the  
following tissue(s): RPE and Choroid. The library was  
constructed according to Bonaldo, Lennon and Soares,  
Genome Research, 6:791-806, 1996. First strand cDNA  
synthesis was primed with an oligo-dT primer containing a  
Not I site. Double stranded cDNA was ligated to an EcoR I

adaptor, digested with Not I, and cloned directionally  
into p7773-Pac vector. The oligonucleotide used to prime  
the synthesis of first-strand cDNA contains a library tag  
sequence that is located between the Not I site and the  
(dT)<sub>18</sub> tail. The sequence tag for this library is ACCCA.  
This library was created for the program, Gene Discovery  
in the Visual System, supported by National Eye Institute  
(NEI)."

BASE COUNT 116 a 157 c 111 g 109 t  
ORIGIN

Query Match 89.2%; Score 21.4; DB 14; Length 493;  
Best Local Similarity 95.7%; Pred. No. 27;  
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
QY 1 GCTTCTTTGCCAAGAGCGCGC 23  
DB 8 GCGTCTTTGCCAAGAGCGCGC 30

Search completed: March 17, 2003, 13:09:15  
Job time : 851.387 secs

GenCore version 5.1.4-P5-4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 09:43:42 ; Search time 202.495 Seconds

(without alignments)  
3161.870 Million cell updates/sec

Title: US-09-836-439-4

Sequence: 1 aagaaatactagacaagca 22

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2054640 segs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenBank1:\*

1: gb-ba:\*

2: gb-hlg:\*

3: gb-in:\*

4: gb-om:\*

5: gb-ov:\*

6: gb-pat:\*

7: gb-ph:\*

8: gb-pl:\*

9: gb-pr:\*

10: gb-ro:\*

11: gb-sts:\*

12: gb-sy:\*

13: gb-un:\*

14: gb-vi:\*

15: em-ba:\*

16: em-fun:\*

17: em-hum:\*

18: em-in:\*

19: em-mu:\*

20: em-om:\*

21: em-or:\*

22: em-ov:\*

23: em-pat:\*

24: em-ph:\*

25: em-pl:\*

26: em-ro:\*

27: em-sts:\*

28: em-un:\*

29: em-vi:\*

30: em-hlg-hum:\*

31: em-hlg-inv:\*

32: em-hlg-mus:\*

33: em-hlg-other:\*

34: em-hlg-pin:\*

35: em-hlg-rod:\*

36: em-hlg-mam:\*

37: em-hlg-vrt:\*

38: em-sy:\*

39: em-higo-hum:\*

40: em-higo-mus:\*

41: em-higo-other:\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20.4	92.7	133095	2	AC046146
2	20.4	92.7	152147	2	AC095531
3	20.4	92.7	172497	2	AC117913
4	20.4	92.7	188795	2	AC113320
5	19.4	88.2	179767	2	AC069223
6	19.4	88.2	180638	9	AC068763
7	19.4	88.2	185841	2	AC027080
8	19.4	88.2	201416	2	AC130437
9	19.4	88.2	218859	2	AC022912
10	19.4	88.2	274349	2	AC093623
11	19.4	88.2	113940	2	AF005572
12	18.8	85.5	2052	5	AF012746
13	18.8	85.5	2209	8	SCYPL252C
14	18.8	85.5	2411	8	SCYPL253C
15	18.8	85.5	37808	8	SC38RCXYI
16	18.8	85.5	41578	3	CBR646G14
17	18.8	85.5	52495	2	AC109155
18	18.8	85.5	52912	2	AC130512
19	18.8	85.5	63484	2	AC108916
20	18.8	85.5	64417	2	AC131268
21	18.8	85.5	96036	2	AC119622
22	18.8	85.5	97454	9	AL138147
23	18.8	85.5	99374	9	AC092031
24	18.8	85.5	112632	2	AL137158
25	18.8	85.5	113955	9	AC068291
26	18.8	85.5	114457	2	AC095684
27	18.8	85.5	120527	2	AP005487
28	18.8	85.5	132072	9	AC004841
29	18.8	85.5	140656	5	AF112374
30	18.8	85.5	141138	2	AL845428
31	18.8	85.5	143496	2	AC111035
32	18.8	85.5	143823	9	AC019195
33	18.8	85.5	151040	2	AC068389
34	18.8	85.5	152161	9	AL359074
35	18.8	85.5	154348	2	AC069480
36	18.8	85.5	156195	9	AC093799
37	18.8	85.5	157038	2	AC099275
38	18.8	85.5	157823	2	AL161795
39	18.8	85.5	158075	2	AC123076
40	18.8	85.5	159386	2	AC098370
41	18.8	85.5	163830	2	AC068235
42	18.8	85.5	166292	2	CNS080C8
43	18.8	85.5	16764	2	AC115875
44	18.8	85.5	168199	9	CNS01DYS
45	18.8	85.5	169295	9	AL732602

## ALIGNMENTS

RESULT 1  
AC046146 133095 bp DNA 11near HTG 16-OCT-2001  
LOCUS  
DEFINITION Mus musculus chromosome 12 clone RP23-321N21, \*\*\* SEQUENCING IN  
ACCESSION AC046146  
VERSION AC046146.6 GI:16118085  
KEYWORDS HTG; HTGS\_PHASE1.  
SOURCE Mus musculus  
ORGANISM Mus musculus

REFERENCE  
1 (bases 1 to 133095)  
Mammalia: Eutheria: Rodentia: Sciurognathi: Muridae: Murinae; Mus.  
Metzker, M.L., Lewis, L.R., Hume, J., Edwards, C., Harris, C.,  
Dederich, D., Thomas, S., Okwuonu, G., Carlock, C., Garner, T.,

TITLE  
JOURNAL  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

Addison, S., Pace, A., Williams, G., Bonnin, D., Brooks, A., Brown, J.,  
Bahay, C., Bunac, C., Burkett, C., Chacko, J., Chen, G., Chen, Z.,  
Cox, C., Davis, C., Delgado, O., Ding, Y., Dugan-Rocha, S.,  
Fernandez, C., Ferraguto, J., Forcum-Tansey, J., Gill, R.,  
Gorrell, J. H., Gunaratne, P., Haller, G., Hernandez, J., Hogues, M.,  
Hosack, H., Hou, X., Huber, J., Jackson, L., Jia, Y., Kelly, J., Kelly, S.,  
Kovar, C., Liu, D., Liu, W., Louised, H., Lozano, R. J., Martin, R.,  
Massey, E., McLeod, M. P., Mei, G., Moore, S., Morgan, M., Morris, S.,  
Neal, D., Nelson, A., Nguyen, R., Nguyen, N., Ogih, M., Parish, B.,  
Perez, L., Reiter, D., Say, J., Shen, H., Vasquez, L., Wallington, S.,  
Williamson, A., Wrensford, G., Zhou, X., Bouck, J., Hodgson, A.,  
Muzny, D. M., Rives, M., Scherer, S., Sodergren, E., Weinstein, G.,  
Worley, K., and Gibbs, R.

Unpublished  
2 (bases 1 to 133095)  
Direct Submission

Submitted (13-APR-2000) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA  
On Oct 14, 2001 this sequence version replaced g1:11094634.

----- Genome Center  
Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: [hgsc.helpebcm.tmc.edu](mailto:hgsc.helpebcm.tmc.edu)  
----- Project Information  
Center Project name: MADY  
Center clone name: RP23-321N21  
----- Summary Statistics  
Sequencing vector: M13; L08E21  
Chemistry: Dye-terminator Big Dye; 94% of reads  
Assembly program: Phrap; version 0.990329  
Consensus quality: 118378 bases at least Q40  
Consensus quality: 144876 bases at least Q30  
Consensus quality: 155160 bases at least Q20  
Estimated insert size: 148882; sum-of-contigs estimation  
Quality coverage: 0x in Q20 bases; agarose-ff estimation  
Quality coverage: 2.1x in Q20 bases; sum-of-contigs estimation

----- NOTE: Estimated insert size may differ from sequence length  
(see <http://www.hgsc.bcm.tmc.edu/docs/genbank.draft.data.html>).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 31 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1 10336: contig of 10336 bp in length  
\* 10337 10436: gap of unknown length  
\* 10437 19208: contig of 8772 bp in length  
\* 19209 19308: gap of unknown length  
\* 19309 26724: contig of 7416 bp in length  
\* 26725 26824: gap of unknown length  
\* 26825 33512: contig of 6688 bp in length  
\* 33513 33612: gap of unknown length  
\* 33613 33888: contig of 5776 bp in length  
\* 33889 39489: gap of unknown length  
\* 39490 45330: contig of 5842 bp in length  
\* 45331 50898: gap of unknown length  
\* 50899 50998: gap of unknown length  
\* 50999 56466: gap of unknown length  
\* 56467 62096: contig of 5530 bp in length  
\* 62097 62196: gap of unknown length  
\* 62197 65971: contig of 3775 bp in length  
\* 65972 66071: gap of unknown length  
\* 66072 70109: contig of 4038 bp in length

70110 70209: gap of unknown length  
\* 70210 72595: contig of 2286 bp in length  
\* 72596 74972: gap of unknown length  
\* 74973 75072: contig of 2277 bp in length  
\* 75073 79255: gap of unknown length  
\* 79256 79355: contig of 4183 bp in length  
\* 79356 83184: contig of 3828 bp in length  
\* 83184 83283: gap of unknown length  
\* 83284 86781: contig of 3498 bp in length  
\* 86782 90021: contig of 3140 bp in length  
\* 90022 90121: gap of unknown length  
\* 90122 93597: contig of 3476 bp in length  
\* 93598 93697: gap of unknown length  
\* 93698 97229: contig of 3552 bp in length  
\* 97230 97329: gap of unknown length  
\* 97330 100516: contig of 3187 bp in length  
\* 100517 100616: gap of unknown length  
\* 100617 104190: contig of 3574 bp in length  
\* 104191 104290: gap of unknown length  
\* 104291 107955: contig of 3665 bp in length  
\* 107956 108055: gap of unknown length  
\* 108056 111930: contig of 3875 bp in length  
\* 111931 112030: gap of unknown length  
\* 112031 114125: contig of 2095 bp in length  
\* 114126 114225: gap of unknown length  
\* 114226 116520: contig of 2295 bp in length  
\* 116521 116620: gap of unknown length  
\* 116621 118829: contig of 2209 bp in length  
\* 118830 118930: gap of unknown length  
\* 118931 119299: gap of unknown length  
\* 119300 121281: contig of 2352 bp in length  
\* 121282 121381: gap of unknown length  
\* 121382 123762: contig of 2381 bp in length  
\* 123763 123862: gap of unknown length  
\* 123863 126537: contig of 2674 bp in length  
\* 126537 126636: gap of unknown length  
\* 126637 130483: contig of 3847 bp in length  
\* 130484 130583: gap of unknown length  
\* 130584 130584: contig of 2512 bp in length.

FEATURES  
source  
1. 133095  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/chromosome="12"  
/clone="RP23-321N21"

BASE COUNT 34946 a 30809 c 29300 g 35021 t 3019 others

ORIGIN  
Query Match 92.7%; Score 20.4; DB 2; Length 133095;  
Best Local Similarity 95.5%; Pred. No. 65;  
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 AACAAATCTAGACAAACAA 22  
|||||  
DB 71615 AAGCAAAATCTAGACAAACAA 71636

RESULT 2  
AC095531  
LOCUS  
DEFINITION Rattus norvegicus clone CH230-8N2, \*\*\* SEQUENCING IN PROGRESS \*\*\*  
AC095531 152147 bp DNA linear HTG 10-JUL-2002  
58 unordered pieces.  
AC095531  
AC095531.3 GI:21716992  
VERSION  
HTG: HTGS\_PHASE1.  
SOURCE Norway rat.  
ORGANISM Rattus norvegicus  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
Rattus.  
1 (bases 1 to 152147)  
Muzny, D. M., Adams, C., Adio-Oduola, B., Ali-osman, F. R., Allen, C.,



Alsbrooks, S.L., Amaralunge, H.C., Are, J.R., Ayele, M., Banks, T.,  
 Barbra, J., Benton, J., Blimage, K., Blankenburg, K., Bonnin, D.,  
 Bouck, J., Bowe, S., Brileva, M., Brown, E., Brown, M., Bryant, N.P.,  
 Buhay, C., Burch, P., Burkett, C., Burrell, K.L., Byrd, N.C.,  
 Cartron, T.F., Carter, M., Cavazos, S.R., Chacko, J., Chavez, D.,  
 Chen, G., Chen, R., Chen, Z., Chowdhry, I., Christopoulos, C.,  
 Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R.,  
 Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A.,  
 Delaney, K.R., Delgado, O., Dem, A.L., Ding, Y., Dink, H.H.,  
 Douthwaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J.,  
 Earnhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Escoto, M.,  
 Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P.,  
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 Garrell, J.H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K.,  
 Harris, C., Harris, K., Hart, M., Havlek, P., Hawes, A., Hernandez, J.,  
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 Peters, L., Pickens, R., Primus, E., Pu, L.L., Quiles, M., Ren, Y.,  
 Rivers, M., Rojas, A., Rojubokan, I., Rolfe, M., Ruiz, S., Savary, G.,  
 Scherer, S., Scott, G., Shen, H., Shoohtari, N., Sisson, I.,  
 Sodergren, E., Sonaike, T., Sparks, A., Stanley, H., Stone, H.,  
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 Usmani, K., Vasquez, L., Vera, Y., Villalon, D., Vinson, R., Wang, Q.,  
 Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S.,  
 Williams, G., Williamson, A., Wlecczyk, R., Woodson, S., Worley, K.,  
 Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Nelson, D.,  
 Wulfsberg, G. and Gibbs, R.

# Direct Submission

2 (bases 1 to 152147)

Worley, K.C.

Direct Submission

Submitted (17-SEP-2001) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 152147)

Worley, K.C.

Direct Submission

Submitted (10-JUL-2002) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

On Jul 9, 2002 this sequence version replaced gi:17942049.

Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

Project Information

Center project name: GCPY

Center clone name: CH230-BN2

Summary Statistics

Sequencing vector: Plasmid

Chemistry: Dye-terminator Big Dye: 100% of reads

Assembly program: Phrap: version 0.990329

Consensus quality: 92864 bases at least Q40

Consensus quality: 97631 bases at least Q30

Consensus quality: 101839 bases at least Q20

NOTE: Estimated insert size may differ from sequence length

(see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_dir/c\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_dir/c_data.html)).

NOTE: This is a 'working draft' sequence. It currently

\* consists of 58 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

1	1151: contig of 1151 bp in length
1152	gap of unknown length
1252	contig of 1454 bp in length
2705	gap of unknown length
2805	contig of 1516 bp in length
4321	gap of unknown length
4421	contig of 1438 bp in length
5859	gap of unknown length
5955	contig of 1267 bp in length
5960	gap of unknown length
7225	contig of 1117 bp in length
7325	gap of unknown length
8443	contig of 1160 bp in length
8544	gap of unknown length
9703	contig of 1160 bp in length
9704	gap of unknown length
9804	contig of 1232 bp in length
11036	gap of unknown length
11135	contig of 1788 bp in length
12923	gap of unknown length
13023	contig of 1152 bp in length
14175	gap of unknown length
14275	contig of 1937 bp in length
16212	gap of unknown length
16312	contig of 1579 bp in length
17891	gap of unknown length
17991	contig of 1341 bp in length
19332	gap of unknown length
19433	contig of 1413 bp in length
20845	gap of unknown length
20945	contig of 1838 bp in length
22783	gap of unknown length
22883	contig of 1036 bp in length
23919	gap of unknown length
24019	contig of 1169 bp in length
25188	gap of unknown length
25288	contig of 1146 bp in length
26434	gap of unknown length
26534	contig of 1405 bp in length
27939	gap of unknown length
28039	contig of 1781 bp in length
29820	gap of unknown length
29920	contig of 1489 bp in length
31409	gap of unknown length
31509	contig of 1798 bp in length
33307	gap of unknown length
33407	contig of 2019 bp in length
35425	gap of unknown length
35525	contig of 1354 bp in length
36880	gap of unknown length
36980	contig of 1549 bp in length
38529	gap of unknown length
38629	contig of 1517 bp in length
40145	gap of unknown length
40245	contig of 2271 bp in length
42517	gap of unknown length
42617	contig of 1675 bp in length
44292	gap of unknown length
44392	contig of 2316 bp in length
46708	gap of unknown length
46808	contig of 2434 bp in length
49242	gap of unknown length
49342	contig of 1916 bp in length
51258	gap of unknown length
51358	contig of 2002 bp in length
53360	gap of unknown length
53460	contig of 1688 bp in length
55148	gap of unknown length
55248	contig of 1688 bp in length

```

* 55249 58677: contig of 3429 bp in length
* 58678 58777: gap of unknown length
* 58778 60637: contig of 1860 bp in length
* 60638 60737: gap of unknown length
* 60738 63930: contig of 3193 bp in length
* 64031 66398: contig of 2368 bp in length
* 66399 66498: gap of unknown length
* 66499 69978: contig of 3480 bp in length
* 69979 70078: gap of unknown length
* 70079 72552: contig of 2474 bp in length
* 72553 74752: gap of unknown length
* 74753 74852: contig of 2100 bp in length
* 74853 77949: contig of 3097 bp in length
* 77950 78050: gap of unknown length
* 78050 80099: contig of 1950 bp in length
* 80100 82883: gap of unknown length
* 82884 82984: contig of 2784 bp in length
* 82985 86241: gap of unknown length
* 86242 86341: contig of 3257 bp in length
* 86342 90145: gap of unknown length
* 90146 90245: contig of 3805 bp in length
* 90246 92780: gap of unknown length
* 92781 92881: contig of 2535 bp in length
* 92882 97022: gap of unknown length
* 97023 97123: contig of 4142 bp in length
* 97124 99914: gap of unknown length
* 99915 100014: contig of 2792 bp in length
* 100015 103193: gap of unknown length
* 103194 103292: contig of 3178 bp in length
* 103293 107022: gap of unknown length
* 107023 107122: contig of 3730 bp in length
* 107123 110597: gap of unknown length
* 110598 110697: contig of 3475 bp in length
* 110698 116077: gap of unknown length
* 116078 116178: contig of 5380 bp in length
* 116179 120578: gap of unknown length
* 120579 120678: contig of 4401 bp in length
* 120679 125200: gap of unknown length

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```

Query Match      92.7%  Score 20.4;  DB 2;  Length 152147;
Best Local Similarity 95.5%  Pred. No. 63;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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OY 1 AAGAAAAATCTAGACAGCAA 22
Db 141943 AAGAAAAATGTAGACAGCAA 141964

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RESULT 3
AC117913/c 172497 bp DNA linear HTG 18-JUL-2002
LOCUS Rattus norvegicus clone CH230-35D24, *** SEQUENCING IN PROGRESS
DEFINITION *** 59 unordered pieces.
ACCESSION AC117913
VERSION AC117913.4 GI:21747159
KEYWORDS HTG; HTGS_PHASE1.
SOURCE Norway rat.
ORGANISM Rattus norvegicus
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 172497)
Munzay,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,
Alshbrooks,S.L., Amaralunga,H.C., Are,J.R., Ayele,M., Banks,T.,
Barbata,J., Benton,J., Bimaga,K., Blankenburg,K., Bonnin,D.,
Bouck,J., Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P.,
Bunay,C., Butch,P., Burkett,C., Burrell,K.L., Byrd,N.C.,
Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D.,
Chen,G., Chen,R., Chen,Z., Chowdhry,I., Christopoulos,C.,
Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R.,

```

## COMMENT

```

REFERENCE
AUTHORS
TITLE
JOURNAL

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```

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

```

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Davila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A.,
Delaney,K.R., Delgado,O., Dena,A.L., Ding,Y., Dinh,H.H.,
Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Durbin,K.J.,
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Peters,L., Pickens,R., Primus,E., Pu,L.L., Quiles,M., Ren,Y.,
Rives,M., Rojas,A., Rojokan,I., Rolfe,M., Ruiz,S., Savery,G.,
Scherer,S., Scott,G., Shen,H., Shooshari,N., Sisson,I.,
Sodergren,E., Sonalke,T., Sparks,A., Stanley,H., Stone,H.,
Sutton,A., Svatek,A., Taber,P., Tamerisa,A., Tamerisa,K., Tang,H.,
Tansley,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S.,
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Wang,S., Ward-Moore,S., Warren,M., Washington,C., Watlington,S.,
Williams,G., Williamson,A., Wleczek,R., Wooden,S., Worley,K.,
Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,
Weinstock,G., and Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 172497)
Worley,K.C.
Direct Submission
Submitted (11-APR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 172497)
Worley,K.C.
Direct Submission
Submitted (18-JUL-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Jul 14, 2002 this sequence version replaced gi:20258282.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GYPM
Center clone name: CH230-35D24
----- Summary Statistics
Sequencing vector: plasmid;
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 118901 bases at least Q40
Consensus quality: 125278 bases at least Q30
Consensus quality: 129335 bases at least Q20
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* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_data.html).
* NOTE: This is a "working draft" sequence. It currently
* consists of 59 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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* 1 1133: contig of 1133 bp in length
* 1134 1233: gap of unknown length
* 1234 2457: contig of 1224 bp in length
* 2458 2557: gap of unknown length
* 2558 3893: contig of 1336 bp in length
* 3894 3994: gap of unknown length
* 3994 5079: contig of 1086 bp in length
* 5080 5179: gap of unknown length
* 5180 6739: contig of 1560 bp in length
* 6740 8037: gap of unknown length
* 8038 8137: gap of unknown length
* 8138 9769: contig of 1632 bp in length
* 9770 9869: gap of unknown length
* 9870 11173: contig of 1304 bp in length
* 11174 11273: gap of unknown length
* 11274 12323: contig of 1056 bp in length
* 12330 12429: gap of unknown length
* 12430 14363: contig of 1934 bp in length
* 14364 14463: gap of unknown length
* 14464 15857: contig of 1394 bp in length
* 15858 15957: gap of unknown length
* 15958 17164: contig of 1207 bp in length
* 17165 17264: gap of unknown length
* 17265 18407: contig of 1143 bp in length
* 18408 18508: gap of unknown length
* 18509 20371: contig of 1864 bp in length
* 20372 20471: gap of unknown length
* 20472 21793: contig of 1321 bp in length
* 21793 21892: gap of unknown length
* 21893 23539: contig of 1647 bp in length
* 23540 23639: gap of unknown length
* 23640 24789: contig of 1150 bp in length
* 24790 24889: gap of unknown length
* 24890 26903: contig of 2014 bp in length
* 26904 27003: gap of unknown length
* 27004 28488: contig of 1485 bp in length
* 28489 28588: gap of unknown length
* 28589 29829: contig of 1241 bp in length
* 29830 29929: gap of unknown length
* 29930 31463: contig of 1533 bp in length
* 31463 31562: gap of unknown length
* 31563 33655: contig of 2093 bp in length
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* 33756 35134: contig of 1379 bp in length
* 35135 35234: gap of unknown length
* 35235 37277: contig of 2043 bp in length
* 37278 37377: gap of unknown length
* 37378 38782: contig of 1405 bp in length
* 38783 38882: gap of unknown length
* 38883 40412: contig of 1530 bp in length
* 40413 40512: gap of unknown length
* 40513 42065: contig of 1553 bp in length
* 42066 42165: gap of unknown length
* 42166 44423: contig of 2258 bp in length
* 44424 44523: gap of unknown length
* 44524 45924: contig of 1401 bp in length
* 45925 46024: gap of unknown length
* 46025 47783: contig of 1755 bp in length
* 47784 47883: gap of unknown length
* 47884 50262: contig of 2379 bp in length
* 50263 50362: gap of unknown length
* 50363 52442: contig of 2080 bp in length
* 52443 52542: gap of unknown length
* 52543 55030: contig of 2488 bp in length
* 55031 55130: gap of unknown length
* 55131 57183: contig of 2453 bp in length
* 57184 57683: gap of unknown length
* 57684 59735: contig of 2052 bp in length
* 59736 59835: gap of unknown length
* 59836 62023: contig of 2188 bp in length
* 62024 62123: gap of unknown length
* 62124 65327: contig of 3204 bp in length

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* 65328 65427: gap of unknown length
* 65428 67771: contig of 2344 bp in length
* 67772 67871: gap of unknown length
* 67872 70986: contig of 3115 bp in length
* 70987 71086: gap of unknown length
* 71087 74496: contig of 3410 bp in length
* 74497 74596: gap of unknown length
* 74597 76391: contig of 1795 bp in length
* 76392 76491: gap of unknown length
* 76492 80570: contig of 4079 bp in length
* 80571 80671: gap of unknown length
* 80672 84667: contig of 3997 bp in length
* 84668 84767: gap of unknown length
* 84768 88261: contig of 3494 bp in length
* 88262 88361: gap of unknown length
* 88362 91664: contig of 3303 bp in length
* 91665 91764: gap of unknown length
* 91765 96059: contig of 4295 bp in length
* 96060 96159: gap of unknown length
* 96160 101392: contig of 5233 bp in length
* 101393 101492: gap of unknown length
* 101493 104583: contig of 3091 bp in length
* 104584 104683: gap of unknown length
* 104684 109409: contig of 4726 bp in length
* 109410 109510: gap of unknown length
* 109511 114694: contig of 5185 bp in length
* 114695 114794: gap of unknown length
* 114795 119788: contig of 4994 bp in length
* 119789 119888: gap of unknown length
* 119889 126089: contig of 6200 bp in length
* 126089 126188: gap of unknown length
* 126189 134246: contig of 8058 bp in length
* 134247 134346: gap of unknown length
* 134347 140261: contig of 5915 bp in length

Query Match
Best Local Similarity 92.7%; Score 20.4; DB 2; Length 172497;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAGAAAATCTAGACAGCAA 22
Db 45058 AGAGAAAATGTAGACAGCAA 45037

RESULT 4
AC113320
LOCUS 188795 bp DNA linear HTG 24-AUG-2002
DEFINITION Mus musculus clone RP23-445E16, WORKING DRAFT SEQUENCE, 11 ordered
pieces.
AC113320
AC113320.2 GI:22474991
VERSION HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_FULLTOP.
KEYWORDS house mouse.
SOURCE Mus musculus
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 188795)
AUTHORS Birren,B., Nusbaum,C. and Lander,E.
JOURNAL Mus musculus, clone RP23-445E16
TITLE Unpublished
REFERENCE 2 (bases 1 to 188795)
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N.,
Anderson,S., Barina,N., Bastien,V., Boguslavsky,L., Boukhalter,B.,
Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
Choepe,Y., Collangelo,M., Collins,S., Collymore,A., Cook,A.,
Cooke,P., DeArrellano,K., Dewar,K., Diaz,J.S., Dodge,S., Fato,S.,
Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
Kamali,A., Karatas,A., Kells,C., Lacombe,K., Lamazares,R.,
Lander,T., Lehoczy,J., Levine,R., Liu,G., MacLean,C.,
Macdonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M.,
McEwan,P., McKernan,K., Meldrum,J., Meneus,L., Milhova,T.,

```

TITLE  
JOURNAL  
REFERENCE  
AUTHORS

Submitted (28-FEB-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
(bases 1 to 188795)  
Britten, B., Nishikawa, C., Lander, E., Ali, A., Allen, N., Anderson, S., Barina, N., Bastien, Y., Bloom, T., Boguslavsky, L., Boukhgalter, B., Camarata, J., Chang, J., Chazaro, B., Choquet, Y., Collymore, A., Cook, A., Cooke, P., DeArrellano, K., Dewar, K., Diaz, J.S., Dodge, S., Fato, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardina, S., Gird, S., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, G., Maclean, C., Macdonald, P., Major, J., Matthews, C., McCarthy, M., Meldrum, J., Menes, L., Mihova, T., Mieng, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Raymond, C., Rella, R., Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schupack, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE  
JOURNAL  
COMMENT

Submitted (24-AUG-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
On Aug 24, 2002 this sequence version replaced g1:16997726.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1997)  
http://ftp.genome.washington.edu/RM/RepeatMasker.html

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: W1BR

Web site: http://www-seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu

Project Information

Center project name: L22784

Center clone name: 445\_E\_16

Summary Statistics

Sequencing vector: Plasmid; n/a; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 185600 bases at least Q40

Consensus quality: 186954 bases at least Q30

Consensus quality: 187365 bases at least Q20

Insert size: 18800; agarose-IP

Insert size: 187795; sum-of-coverage

Quality coverage: 9.1 in Q20 bases; agarose-IP

Quality coverage: 9.1 in Q20 bases; sum-of-coverage

NOTE: This is a 'working draft' sequence. It currently consists of 11 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have been provided by the submitter.

This sequence will be replaced

by the finished sequence as soon as it is available and

the accession number will be preserved.

1 47353: contig of 47353 bp in length

47354 47453: gap of 100 bp

47454 48767: contig of 1314 bp in length

48768 48867: gap of 100 bp

48868 50642: contig of 1775 bp in length

# FEATURES

## source

50643 50742: gap of 100 bp  
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52086 54528: contig of 2443 bp in length  
54529 54628: gap of 100 bp  
54629 58116: contig of 3488 bp in length  
58117 58216: gap of 100 bp  
58217 60812: contig of 2596 bp in length  
60813 60912: gap of 100 bp  
60913 62765: contig of 1853 bp in length  
62766 62865: gap of 100 bp  
62866 72529: contig of 9664 bp in length  
72530 72629: gap of 100 bp  
72630 96147: contig of 23518 bp in length  
96148 96247: gap of 100 bp  
96248 188795: contig of 92548 bp in length.

Location/Qualifiers

1. 188795

/organism="Mus musculus"

/db\_xref="taxon:10090"

/clone="RP23-445E16"

/clone\_11b="RP23-Female Mouse BAC"

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/note="assembly-fragment"

clone\_end:SP6

vector\_side:left"

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/note="assembly-fragment"

48868 50642

/note="assembly-fragment"

50743 51985

/note="assembly-fragment"

52086 54528

/note="assembly-fragment"

54629 58116

/note="assembly-fragment"

58217 60812

/note="assembly-fragment"

60913 62765

/note="assembly-fragment"

62866 72529

/note="assembly-fragment"

72630 96147

/note="assembly-fragment"

96248 188795

/note="assembly-fragment"

/note="assembly-fragment"

vector\_side:right"

BASE COUNT 57294 a 37412 c 36369 g 56709 t 1011 others

# ORIGIN

## Query Match

Best Local Similarity 95.5%; Pred. No. 61;

Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Y 1 AAGAAAAAATCTAGACAGCAA 22

Db 36223 AAGAAAAAATCTAGACAGCAA 36244

# RESULT 5

## LOCUS

AC069223 179767 bp DNA linear PRI 30-MAR-2001

## DEFINITION

Homo sapiens 3 BAC RP11-398021 (Roswell Park Cancer Institute Human BAC Library) complete sequence.

## ACCESSION

AC069223 GI:13489121

## KEYWORDS

## SOURCE

## ORGANISM

Homo sapiens.

Homo sapiens.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 179767)

## AUTHORS

AUTHORS	TITLE	JOURNAL	REFERENCE	AUTHORS	TITLE	JOURNAL	REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
Auzuny D.M., Adams C., Adio-odunla B., Al-ozman F.R., Allen C., Alsbrooks S.L., Amaratunga H.C., Are J.R., Banks T., Barbataia J., Benton J., Bimarge K., Blankenburg K., Bonnin D., Bouck J., Bowle S., Brieva M., Brown E., Brown M., Bryant N.P., Buhay C., Butch P., Burkett C., Burrell K.L., Byrd N.C., Caron T.F., Carter M., Cavazos S.R., Chacko J., Chavez D., Chen G., Chen R., Chen Z., Chowdhry I., Christopoulos C., Cleveland C.D., Cox C., Coyle M.D., Dathorne S.R., David R., Davila M.L., Davis C., Day-Carroll L., Dederich D.A., Delaney K.R., Delgado H., Den A.L., Ding Y., Dinh H.H., Douthwaite K.J., Draper H., Dugan-Rocha S., Durbin K.J., Earnhart C., Edger D., Edwards C.C., Elhaj C., Escotto M., Falls T., Fernigato D., Flagg N., Ford J., Foster P., Frantz P., Gabisi A., Gao J., Garcia A., Garner T., Garza N., Gill R., Gorrell J.H., Guevara W., Gunaratne P., Hale S., Hamilton K., Harris C., Harris K., Hart M., Hayla P., Hawes A., He X., Hernandez J., Hernandez O., Hodgson A., Hogues M., Holloway C., Hollins B., Homsl F., Howard S., Huber J., Hulyk S., Hume J., Jackson L.E., Jacobson B., Jia Y., Johnson R., Jolivet S., Joudan S., Karlsson E., Kelly S., Khan U., King L., Korvan J., Kover C., Klatov C.J., Kureshi A., Landry N., Leal B., Lewis L.C., Lewis L., Li J., Li Z., Lichtarge O., Liew C., Liu J., Liu W., Lounsgaard H., Lozano R.J., Lu X., Lucier R., Lucier R., Luna R., Ma J., Maheshwari M., Mapa P., Martin R., Matindale A., Martinez E., Massey E., Maxwell E., McLeod M.P., Meador M., Mel G., Metcker M., Miner G., Miner Z., Mitchell T., Mohabbat K., Moore S., Morgan M., Moorish T., Morris S., Moser M., Neal D., Nelson D., Newton J., Newton N., Nguyen A., Nguyen N., Nguyen N., Nickerson E., Nwokweto S., Ogun M., Okunolu G., Orgunye N., Oyedro R., Pace A., Payton B., Peary J., Perez L., Peters L., Pickens R., Pitmus E., Pul L., Quiles M., Ren Y., Rivers M., Rojas A., Rojibokan I., Rolfe M., Ruiz S., Savery G., Scherer S., Scott G., Shen H., Shoshitari N., Sisson I., Sodergren E., Sonalke T., Sparks A., Stanley H., Stone H., Sutton A., Swalek A., Tabot P., Tamerisa A., Tamerisa K., Tang H., Tansey J., Taylor C., Taylor T., Telford B., Thomas N., Thomas S., Usmani K., Vera V., Villalón D., Vanson R., Wall R., Wang S., Ward-Moore S., Warren R., Washington C., Watlington S., Williams G., Williamson A., Wleczyk R., Woodson S., Worley K., Wu C., Wu Y., Wu Y.F., Zhou J., Zorrilla S., Naylor S.L. and Gibbs R.	Direct Submision	2 (bases 1 to 179767)	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	On Mar 30, 2001 this sequence version replaced gi:13430920.
Worley K.C.	Direct Submision	3 (bases 1 to 179767)	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	On Mar 30, 2001 this sequence version replaced gi:13430920.
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Worley K.C.	Direct Submision	3 (bases 1 to 179767)	Submitted (30-MAR-2001)	Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA	Submitted (30-MAR-2001)						

flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: this sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://gc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

## QUALSTAT-REPORT-----

```
----- Summary Statistics
Contig length:
```

Count: length:	179767
Phrap values in estimate:	179551
Average error rate (BCM-Phrap estimate):	4.4907e-06
Fraction of Phrap values less than 40 :	0.00499023
Number of N's in consensus :	4
	0

Position	Consensus changing edits	Original+Context
1	1	1
2	1	1
3	1	1
4	1	1
5	1	1
6	1	1
7	1	1
8	1	1
9	1	1
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91	1	1
92	1	1
93	1	1
94	1	1
95	1	1
96	1	1
97	1	1
98	1	1
99	1	1
100	1	1

Position	Original+Context	Edited+Context
3	atccaanaa(n)tcgacatc	atccgggaa(v)tcgacatc
107934	agccctctt(n)nacagagact	agccctctt(v)tcagagagact
107935	gccctcttn(n)acagagact	gccctctt(v)tcagagagact
159144	atgccaatca(n)caacactgt	atgccaatca(c)caacactgt

----- Distribution of Quality < 40 Bases

A scatter plot showing the relationship between the number of bases (y-axis) and the Phrap Value Range (x-axis). The y-axis ranges from 0 to 1000 in increments of 100. The x-axis ranges from 5 to 40 in increments of 5. A dashed horizontal line is drawn at y = 500. Data points are represented by asterisks (\*).

Phrap Value Range	Number of bases
5	~100
10	~150
15	~200
20	~250
25	~300
30	~350
35	~400
40	~450

## FEATURES

```

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source    Location/Qualifiers
          1..179767
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           /chromosome="3"
           /clone="RP11-398021"
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           /rpt_family="MIR"
           1202..1410
           /rpt_family="AluYb"
           complement(4054..4428)
           /rpt_family="L2"
           complement(4951..5563)

```



Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as low coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://gc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

## QUALSTAT-REPORT-----

```
----- Summary Statistics -----
Contig length: 180638
Phrap values in estimate: 179362
Average error rate (BCM-Phrap estimate): 1.00944e-05
Fraction of Phrap values less than 40 : 0.00994637
Number of consensus changing edits: 49
Number of N's in consensus : 0
```

```
----- Consensus changing edits -----
Position Original+Context Edited+Context
17175 tgttataga(n)aaagagctc tgttataga(a)aaagagctc
17467 tatagataga(n)acgtgcgtc tatagataga(a)acgtgcgtc
18821 atctagaaga(n)atgataaat atctagaaga(a)atgataaat
18822 tctagaaga(n)tgataaat tctagaaga(a)tgataaat
29835 gatgctta(n)gatgttta gatgctta(g)gatgttta
49503 actcagata(n)acgtgcagc actcagata(a)acgtgcagc
51340 tctcattat(n)gatactcagc tctcattat(a)gatactcagc
52598 ggaataaat(n)catgtacaa ggaataaat(a)catgtacaa
60642 acttaaaaat(n)aaatttcgg acttaaaaat(t)aaatttcgg
60920 caacaagagc(n)taactcatt caacaagagc(g)taactcatt
60921 aacaagagct(n)taactcatt aacaagagct(a)taactcatt
60922 aagaagagct(n)actcattc aagaagagct(a)actcattc
64475 agatgcgctc(a)ccgacatca agatgcgctc(c)ccgacatca
73510 atctcagctc(n)ctgcacatc atctcagctc(a)ctgcacatc
73543 tcaagagat(n)tcctgcctca tcaagagat(c)tcctgcctca
73544 tctgcctca(n)ctcccaaat tctgcctca(g)ctcccaaat
73566 tcccaataa(n)tcggagttac tcccaataa(g)tcggagttac
73567 tcccaataa(n)tgagttaca tcccaataa(c)tgagttaca
73704 ccacatgctc(n)gctaatctt ccacatgctc(t)gctaatctt
73721 ggttgagat(n)acagcgatga ggttgagat(t)acagcgatga
73722 gtagcgacac(n)agctgcgctc gtagcgacac(g)agctgcgctc
73748 tgaacacac(n)tgctgcctc tgaacacac(t)tgctgcctc
73750 tctctatta(n)tcgttcctc tctctatta(c)tcgttcctc
73756 accagctcct(n)gattataga accagctcct(c)gattataga
101317 tttttttt(n)ttaaactca tttttttt(t)ttaaactca
104887 tttttttt(g)gagagagctc tttttttt(t)gagagagctc
109869 gtaggagaga(n)gagagagctc gtaggagaga(g)gagagagctc
110982 gtaggagaga(n)gagagagctc gtaggagaga(g)gagagagctc
113503 gtaggagaga(n)gagagagctc gtaggagaga(g)gagagagctc
113504 gtaggagaga(n)gagagagctc gtaggagaga(g)gagagagctc
125301 tttctccan(n)caatttaac tttctccan(a)caatttaac
128519 aaataacat(n)ataattatc aaataacat(t)ataattatc
13447 ctgataatc(n)gcttttttc ctgataatc(t)gcttttttc
13447 ttcggagat(n)tcctccataa ttcggagat(t)tcctccataa
13442 cagctgctg(n)ctagcctaa cagctgctg(t)ctagcctaa
137769 cctgctag(n)gtggtgctc cctgctag(t)gtggtgctc
```

```
----- Distribution of Quality < 40 Bases -----
1000|
900|
800|
700|
600|
500|
400|
300|
200|
100|
0|
* * * * *
5 10 15 20 25 30 35 40
Phrap Value Range

# bases
1000|
900|
800|
700|
600|
500|
400|
300|
200|
100|
0|
* * * * *
5 10 15 20 25 30 35 40
Phrap Value Range

Version: 1.01 gxf0.
Location/Qualifiers

FEATURES
Query Match 88.2%; Score 19.4; DB 9; Length 180638;
Best Local Similarity 95.2%; Pred. No. 1.6e+02;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 AAGAAAAAATCTGACAGCA 21
Db 163480 AAGAAAAAATCTGACAGCA 163500

RESULT 7
AC027080 185841 bp DNA linear HTG 19-APR-2000
LOCUS Homo sapiens chromosome 3 clone RP11-548023 map 3, WORKING DRAFT
DEFINITION Homo sapiens chromosome 3, clone RP11-548023 map 3, WORKING DRAFT
ACCESSION AC027080.2 GI:7596901
VERSION AC027080
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
1 (bases 1 to 185841)
2 (bases 1 to 185841)
Unpublished
1 (bases 1 to 185841)
2 (bases 1 to 185841)
Biren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
Anderson, S., Baldwin, J., Barna, N., Bastien, V., Beda, F.,
Boguslavsky, L., Bouhassira, B., Brown, A., Burdett, G.,
Campopiano, A., Castle, A., Choquet, Y., Colangelo, M., Collins, S.,
Collins, A., Cooke, P., DeBartolo, K., Dewar, K., Diaz, J.S.,
Dodgson, S., Domino, M., Doyle, M., Ferreira, P., Fitzhugh, W., Gage, D.,
Galagan, J., Gardy, S., Ginde, S., Goyette, M., Graham, L.,
Hendland, J., Hiley, I., Johnson, R., Jones, C., Kahn, L., Karlas, A.,
Klein, J., Lacroque, K., Lamazares, R., Landers, T., Lehoczy, J.,
185841 bp DNA linear HTG 19-APR-2000
Homo sapiens chromosome 3 clone RP11-548023 map 3, WORKING DRAFT
AC027080.2 GI:7596901
HTG; HTGS_PHASE1; HTGS_DRAFT.
Homo sapiens.
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
1 (bases 1 to 185841)
2 (bases 1 to 185841)
Unpublished
1 (bases 1 to 185841)
2 (bases 1 to 185841)
Biren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
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Boguslavsky, L., Bouhassira, B., Brown, A., Burdett, G.,
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Collins, A., Cooke, P., DeBartolo, K., Dewar, K., Diaz, J.S.,
Dodgson, S., Domino, M., Doyle, M., Ferreira, P., Fitzhugh, W., Gage, D.,
Galagan, J., Gardy, S., Ginde, S., Goyette, M., Graham, L.,
Hendland, J., Hiley, I., Johnson, R., Jones, C., Kahn, L., Karlas, A.,
Klein, J., Lacroque, K., Lamazares, R., Landers, T., Lehoczy, J.,
```

# TITLE JOURNAL COMMENT

Levine, R., Lieu, C., Liu, G., Locke, K., MacDonald, P., Margolis, N., McCarthy, M., McEwan, P., McGuirk, A., McKernan, K., McNeeters, R., Melidim, J., Menus, L., Mihova, T., Miranda, C., Mienga, V., Morrow, J., Murphy, T., Naylor, J., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, T.M., Oliver, J., Peterson, K., Plerre, N., Pisanic, C., Pollara, V., Raymond, C., Riley, R., Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J., Teste, S., Theodore, J., Tirrell, A., Travers, M., Triggillo, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G., Zainoun, J., Zimmer, A. and Zody, M.

Submitted (26-MAR-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
On Apr 19, 2000 this sequence version replaced g1:7329443.  
All repeats were identified using RepeatMasker:  
Smith, A.F.A. & Green, P. (1996-1997)  
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

## Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: [sequence-submissions@genome.wi.mit.edu](mailto:sequence-submissions@genome.wi.mit.edu)

## Project Information

Center project name: L8201

Center clone name: 548\_O23

## Summary Statistics

Sequencing vector: M13; M77815; 100% of reads  
Chemistry: Dye-terminator Big Dye; 100% of reads  
Assembly program: Phrap; version 0.960731  
Consensus quality: 172210 bases at least Q40  
Consensus quality: 178990 bases at least Q30  
Consensus quality: 181814 bases at least Q20  
Insert size: 189000; agarose-ef  
Insert size: 183741; sum-of-contents  
Quality coverage: 4.0 in Q20 bases; agarose-ef  
Quality coverage: 4.1 in Q20 bases; sum-of-contents

\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 22 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1 1458: contig of 1458 bp in length  
\* 1459 1558: gap of 100 bp  
\* 1559 3252: contig of 1694 bp in length  
\* 3253 3352: gap of 100 bp  
\* 3353 5248: contig of 1896 bp in length  
\* 5249 5348: gap of 100 bp  
\* 5349 8165: contig of 2817 bp in length  
\* 8166 8265: gap of 100 bp  
\* 8266 11611: contig of 3346 bp in length  
\* 11612 11711: gap of 100 bp  
\* 11712 16128: contig of 4417 bp in length  
\* 16129 16228: gap of 100 bp  
\* 16229 22113: contig of 5885 bp in length  
\* 22114 22213: gap of 100 bp  
\* 22214 28574: contig of 6361 bp in length  
\* 28575 28674: gap of 100 bp  
\* 28675 34052: contig of 5378 bp in length  
\* 34053 34152: gap of 100 bp  
\* 34153 41388: contig of 7236 bp in length  
\* 41389 41488: gap of 100 bp  
\* 41489 50939: contig of 9451 bp in length  
\* 50940 51039: gap of 100 bp  
\* 51040 61135: contig of 10096 bp in length  
\* 61136 61235: gap of 100 bp  
\* 61236 69782: contig of 8547 bp in length  
\* 69783 69882: gap of 100 bp  
\* 69883 77230: contig of 7348 bp in length

## FEATURES source

\* 77231 77330: gap of 100 bp  
\* 77331 86536: contig of 9206 bp in length  
\* 86537 86636: gap of 100 bp  
\* 86637 98162: contig of 11526 bp in length  
\* 98163 98262: gap of 100 bp  
\* 98263 109460: contig of 11198 bp in length  
\* 109461 109560: gap of 100 bp  
\* 109561 121285: contig of 11725 bp in length  
\* 121286 121385: gap of 100 bp  
\* 121386 135542: contig of 14157 bp in length  
\* 135543 135642: gap of 100 bp  
\* 135643 147139: contig of 11497 bp in length  
\* 147140 147239: gap of 100 bp  
\* 147240 161924: contig of 14685 bp in length  
\* 161925 162024: gap of 100 bp  
\* 162025 185841: contig of 23817 bp in length.  
Location/Qualifiers  
1. 185841  
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/db\_xref="taxon:9606"  
/chromosome="3"  
/map="3"  
/clone="RP11-548023"  
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1559. 3252  
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3353. 5248  
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5349. 8165  
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vector\_side:right"  
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86637. 98162  
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98263. 109460  
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109561. 121285  
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147240. 161924  
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162025. 185841  
/note="assembly-fragment"  
BASE COUNT 57091 a 34837 c 33673 g 58130 t 2110 others



ORIGIN

Query Match 88.2%; Score 19.4; DB 2; Length 185841;  
 Best Local Similarity 95.2%; Pred. No. 1.6e+02;  
 Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 AAGAAAAATCTAGACAGCA 21  
 |||||  
 Db 127736 AAGAAAAATCTAGACAGCA 127756

RESULT 8  
 AC130437/c 201416 bp DNA 11linear HTG 10-AUG-2002  
 LOCUS Homo sapiens chromosome 3 clone RP11-740L19, \*\*\* SEQUENCING IN  
 DEFINITION PROGRESS \*\*\*  
 AC130437  
 AC130437.1 GI:22203194  
 KEYWORDS HTG: HTGS\_PHASE1.  
 SOURCE human.  
 ORGANISM Homo sapiens

REFERENCE  
 AUTHORS Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi;  
 Mammalia: Eutheria: Primates: Catarrhini: Hominoidea: Homo.  
 1 (bases 1 to 201416)  
 Muzny, D.M., Adams, C., Adio-Oduola, B., Ali-Osman, F.R., Allen, C.,  
 Alsbrooks, S.L., Amaralunga, H.C., Are, J.R., Ayala, M., Banks, T.,  
 Barbara, J., Benton, J., Blum, K., Blankenburg, K., Bonnin, D.,  
 Bouck, J., Bowe, S., Brileva, M., Brown, E., Brown, M., Bryant, N.P.,  
 Buhay, C., Burch, P., Burkett, C., Burrell, K.L., Byrd, N.C.,  
 Cartron, T.F., Carter, M., Cavazos, S.R., Chacko, J., Chavez, D.,  
 Chen, G., Chen, R., Chen, Z., Chowdhry, I., Christopoulos, C.,  
 Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R.,  
 Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A.,  
 Delaney, K.R., Delgado, O., Dem, A.L., Ding, Y., Dinh, H.H.,  
 Douthett, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J.,  
 Earhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Escotto, M.,  
 Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P.,  
 Gabriel, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R.,  
 Gorrell, J.H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K.,  
 Harris, C., Harris, K., Hart, M., Havlik, P., Hawes, A., Hernandez, J.,  
 Hernandez, O., Hodgson, A., Hughes, M., Holloway, C., Hollins, B.,  
 Homel, F., Howard, S., Huber, J., Hulys, S., Hume, J., Jackson, L.E.,  
 Jacobson, B., Jia, Y., Johnson, R., Jollivet, S., Joudah, S.,  
 Karlsson, E., Kureshi, A., Landry, N., Leal, B., Lewis, L.C., Lewis, L.,  
 Li, J., Li, Z., Lichtarge, O., Lieu, C., Liu, J., Liu, W., Lousaged, H.,  
 Lozada, R.J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J.,  
 Maheshwari, M., Mapua, P., Martin, R., Martindale, A., Martinez, E.,  
 Massey, E., Mawhney, E., McLeod, M.P., Meador, M., Mei, G., Metker, M.,  
 Moser, G., Miner, Z., Mitchell, T., Mohabbat, K., Morgan, M., Morris, S.,  
 Nguyen, N., Nickerson, E., Nwokwuo, S., Ogih, M., Okunodu, G.,  
 Oragunye, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L.,  
 Peters, L., Pickens, R., Primus, E., Pu, L.L., Qules, M., Ren, Y.,  
 Rivers, M., Rojas, A., Rojudoan, I., Rolfe, M., Ruiz, S., Savary, G.,  
 Scherer, S., Scott, G., Shen, H., Shoshitani, N., Sisson, I.,  
 Sodergren, E., Sonalike, T., Sparks, A., Stanley, H., Stone, H.,  
 Sutton, A., Svatok, A., Tabor, P., Tameis, A., Tameis, K., Tang, H.,  
 Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S.,  
 Umani, K., Vasquez, L., Vera, V., Villalón, D., Vinson, R., Wang, Q.,  
 Wang, S., Ward-Moore, S., Warren, R., Washington, C., Wallington, S.,  
 Williams, G., Williamson, A., Wleczek, R., Wooden, S., Worley, K.,  
 Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Nelson, D.,  
 Weinstein, G., and Glbbs, R.

Direct Submission  
 Unpublished  
 2 (bases 1 to 201416)  
 Worley, K.C.  
 Direct Submission  
 Submitted (10-AUG-2002) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA  
 Genom Center

Center: Baylor College of Medicine  
 Center code: BCM  
 Web site: <http://www.hgsc.bcm.tmc.edu/>  
 Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)  
 Project Information  
 Center project name: HDIU  
 Center clone name: RP11-740L19

Summary Statistics  
 Chemistry: Dye-Primer Bodipy: Infinity% of reads  
 Chemistry: Dye-terminator Big Dye: Infinity% of reads  
 Assembly program: Phrap; version 0.990329  
 Consensus quality: 200930 bases at least Q40  
 Consensus quality: 216627 bases at least Q30  
 Consensus quality: 233347 bases at least Q20

NOTE: Estimated insert size may differ from sequence length  
 (see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html)).  
 NOTE: This is a 'working draft' sequence. It currently  
 consists of 10 contigs. The true order of the pieces  
 is not known and their order in this sequence record is  
 arbitrary. Gaps between the contigs are represented as  
 runs of N, but the exact sizes of the gaps are unknown.  
 This record will be updated with the finished sequence  
 as soon as it is available and the accession number will  
 be preserved.

1 2055: contig of 2055 bp in length  
 2056 2155: gap of unknown length  
 2156 4275: contig of 2120 bp in length  
 4276 4375: gap of unknown length  
 4376 9623: contig of 5248 bp in length  
 9624 9723: gap of unknown length  
 9724 26424: contig of 16700 bp in length  
 26424 26522: gap of unknown length  
 26524 42616: contig of 16093 bp in length  
 42617 42716: gap of unknown length  
 42717 59134: contig of 16418 bp in length  
 59135 59234: gap of unknown length  
 59235 77937: contig of 18703 bp in length  
 77938 78037: gap of unknown length  
 78038 98933: contig of 20856 bp in length  
 98934 99033: gap of unknown length  
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Location/Qualifiers  
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 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /chromosome="3"  
 /clone="RP11-740L19"

BASE COUNT 63751 a 38726 c 37375 g 60653 t 911 others

ORIGIN

Query Match 88.2%; Score 19.4; DB 2; Length 201416;  
 Best Local Similarity 95.2%; Pred. No. 1.6e+02;  
 Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 AAGAAAAATCTAGACAGCA 21  
 |||||  
 Db 174985 AAGAAAAATCTAGACAGCA 174965

RESULT 9  
 AC022912 218859 bp DNA 11linear HTG 26-MAY-2000  
 LOCUS Homo sapiens clone RP11-740L19, WORKING DRAFT SEQUENCE, 34  
 DEFINITION UNORDERED PIECES.  
 AC022912  
 AC022912.3 GI:7596818  
 VERSION HTG: HTGS\_PHASE1; HTGS\_DRAFT.  
 KEYWORDS Homo sapiens.  
 SOURCE Homo sapiens  
 ORGANISM Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi;



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Query Match 88.2%: Score 19.4; DB 2: Length 216859;  
Best Local Similarity 95.2%: Pred No. 1.6e+02;  
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTAGACAGCA 21  
|||||  
Db 146022 AAGAAAAATCTAGACAGCA 146042

RESULT 10  
AC093623/c 274349 bp DNA linear HTG 07-SEP-2001  
LOCUS Homo sapiens chromosome UNK clone CTD-2335D13, \*\*\* SEQUENCING IN  
DEFINITION  
AC093623  
PROGRESS \*\*\* 49 unordered pieces.  
AC093623.1 GI:15487445  
VERSION  
KEYWORDS HTG; HTGS\_PHASE1.  
SOURCE  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE  
AUTHORS Waterston, R.H.  
TITLE The sequence of Homo sapiens clone  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 274349)  
AUTHORS Waterston, R.H.

TITLE  
JOURNAL  
COMMENT

Direct Submission  
Submitted (07-SEP-2001) Genome Sequencing Center, Washington  
University School of Medicine, 4444 Forest Park Parkway, St. Louis,  
MO 63108, USA

----- Genome Center -----  
Center: Washington University Genome Sequencing Center  
Center code: WUGSC  
Web site: <http://genome.wustl.edu/gsc/index.shtml>  
----- Project Information -----  
Center project name: H\_MS2335D13  
----- Summary Statistics -----  
Sequencing vector: M13, 1%  
Sequencing vector: plasmid: 99%  
Chemistry: Dye-terminator Big Dye, 99% of reads  
Chemistry: Dye-terminator Big Dye, 99% of reads  
Assembly program: Phrap, version 0.990319  
Consensus quality: 248392 bases at least Q40  
Consensus quality: 257485 bases at least Q30  
Consensus quality: 262193 bases at least Q20  
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\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 49 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1 1390 1389: contig of 1389 bp in length  
\* 1490 1489: gap of unknown length  
\* 2557 2556: contig of 1067 bp in length  
\* 3865 3865: gap of unknown length  
\* 5038 5038: contig of 1073 bp in length  
\* 5139 5138: gap of unknown length  
\* 6763 6762: contig of 1624 bp in length  
\* 8029 8029: gap of unknown length  
\* 8130 8129: gap of unknown length  
\* 9245 9245: contig of 1116 bp in length  
\* 9346 9345: gap of unknown length  
\* 10442 10441: contig of 1096 bp in length  
\* 10542 10541: gap of unknown length  
\* 11789 11788: contig of 1247 bp in length  
\* 11889 11888: gap of unknown length  
\* 13219 13219: contig of 1331 bp in length  
\* 13320 13319: gap of unknown length  
\* 14328 14328: contig of 1009 bp in length  
\* 14329 14428: gap of unknown length  
\* 15578 15578: contig of 1150 bp in length  
\* 15679 15678: gap of unknown length  
\* 17242 17242: contig of 1564 bp in length  
\* 17243 17342: gap of unknown length  
\* 17343 19056: contig of 1714 bp in length  
\* 19057 19156: gap of unknown length  
\* 19157 20355: contig of 1199 bp in length  
\* 20356 20455: gap of unknown length  
\* 20456 21888: contig of 1433 bp in length  
\* 21889 21988: gap of unknown length  
\* 21989 23748: contig of 1760 bp in length  
\* 23749 23848: gap of unknown length  
\* 23849 25791: contig of 1943 bp in length  
\* 25792 25891: gap of unknown length  
\* 25892 29014: contig of 3123 bp in length  
\* 29015 29114: gap of unknown length  
\* 29115 30653: contig of 1539 bp in length  
\* 30654 30753: gap of unknown length  
\* 30754 32579: contig of 1826 bp in length  
\* 32580 32679: gap of unknown length  
\* 32680 34693: contig of 2014 bp in length  
\* 34694 34793: gap of unknown length

```

* 34794 36794: contig of 2001 bp in length
* 36795 36894: gap of unknown length
* 36895 39286: contig of 2392 bp in length
* 39287 39386: gap of unknown length
* 39387 42070: contig of 2684 bp in length
* 42071 42170: gap of unknown length
* 42171 45523: contig of 3353 bp in length
* 45524 45623: gap of unknown length
* 45624 47928: contig of 2305 bp in length
* 47929 48028: gap of unknown length
* 48029 50378: contig of 2350 bp in length
* 50379 50478: gap of unknown length
* 50479 52145: contig of 1667 bp in length
* 52146 52245: gap of unknown length
* 52246 54934: contig of 2689 bp in length
* 54935 55034: gap of unknown length
* 55035 57539: contig of 2505 bp in length
* 57540 57639: gap of unknown length
* 57640 60533: contig of 2394 bp in length
* 60534 60633: gap of unknown length
* 60634 63866: contig of 3323 bp in length
* 63867 63966: gap of unknown length
* 63967 67394: contig of 3428 bp in length
* 67395 67494: gap of unknown length
* 67495 71251: contig of 3757 bp in length
* 71252 71351: gap of unknown length
* 71352 73513: contig of 4162 bp in length
* 73514 75613: gap of unknown length
* 75614 79500: contig of 3887 bp in length
* 79501 79600: gap of unknown length
* 79601 85955: contig of 6355 bp in length
* 85956 86055: gap of unknown length
* 86056 90896: contig of 4841 bp in length
* 90897 90996: gap of unknown length
* 90997 95402: contig of 4406 bp in length
* 95403 95502: gap of unknown length
* 95503 100307: contig of 4805 bp in length
* 100308 100407: gap of unknown length
* 100408 105113: contig of 4706 bp in length
* 105114 105213: gap of unknown length
* 105214 109576: contig of 4363 bp in length
* 109577 109676: gap of unknown length
* 109677 114722: contig of 5046 bp in length
* 114723 114822: gap of unknown length
* 114823 119838: contig of 5016 bp in length
* 119839 119938: gap of unknown length
* 119939 128278: contig of 8340 bp in length
* 128279 128378: gap of unknown length
* 128379 137203: contig of 8825 bp in length
* 137204 137303: gap of unknown length
* 137304 170951: contig of 33648 bp in length
* 170952 171051: gap of unknown length
* 171052 274349: contig of 103298 bp in length.

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## FEATURES

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  /db_xref="taxon:9606"
  /chromosome="UNK"
  /clone="CMD-2335D13"
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1. .1389
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misc_feature
1490. .2556
  /note="assembly_name:Contig22"
misc_feature
2657. .3865
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misc_feature
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  /note="assembly_name:Contig28"
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misc_feature
8130. .9245
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  /note="assembly_name:Contig41"
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misc_feature 20456. .21888
  /note="assembly_name:Contig43"
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  /note="assembly_name:Contig44"
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  /note="assembly_name:Contig46"
misc_feature 29115. .30653
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misc_feature 30754. .32579
  /note="assembly_name:Contig49"
misc_feature 32680. .34693
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misc_feature 34794. .36794
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misc_feature 36895. .39286
  /note="assembly_name:Contig52"
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Query Match 88.2%; Score 19.4; DB 2; Length 274349;
Best Local Similarity 95.2%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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OY 1 AAGAAAAATCTAGACAGCA 21
|||||
DB 220758 AAGAAAAATCTAGACAGCA 220738

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```

RESULT 11
LOCUS AP005572/c 113940 bp DNA linear HTG 24-JUL-2002
DEFINITION Oryza sativa (japonica cultivar-group) chromosome 9 clone
OJ1506_A04, ** SEQUENCING IN PROGRESS ***, in ordered pieces.
ACCESSION AP005572
VERSION AP005572.1 GI:21952940
KEYWORDS HTG; HTGS; PHASE2.
SOURCE Oryza sativa (japonica cultivar-group) (cultivar:Nipponbare) DNA,
clone:OJ1506_A04
ORGANISM Oryza sativa (japonica cultivar-group)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ehrhartoideae; Oryzaeae; Oryza.
REFERENCE 1
AUTHORS Sasaki,T., Matsunoto,T., Hattori,M., Sakaki,Y. and Katayose,Y.
TITLE Oryza sativa nipponbare(GAS) genomic DNA, chromosome 9, BAC
clone:OJ1506_A04
JOURNAL Published Only in Database (2002)
AUTHORS 2 (bases 1 to 113940)
TITLE Direct Submission
AUTHORS Sasaki,T., Matsunoto,T., Hattori,M., Sakaki,Y. and Katayose,Y.
JOURNAL Submitted (23-JUL-2002) Takuji Sasaki, National Institute of
Agricultural Sciences, Rice Genome Research Program, Kannondai
2-1-2, Tsukuba, Ibaraki 305-8602, Japan

```

(E-mail: tsasakienlas.affrc.go.jp, URL: http://rpg.dna.affrc.go.jp/, Tel: 81-298-38-7441, Fax: 81-298-38-7468)  
The nucleotide sequence of this BAC clone was generated by combining Monsanto and RGP-Japan sequencing data.

NOTE: It currently consists of 1 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have provided by the submitter. This sequence will be replaced by the finished sequence as soon as it is available and the accession number will be preserved.

\* NOTE: This is a 'working draft' sequence.  
\* This sequence will be replaced  
\* by the finished sequence as soon as it is available and  
\* the accession number will be preserved.

## FEATURES

source

1. 113940  
Location/Qualifiers  
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/chromosome="9"  
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Best Local Similarity 100.0%; Pred. No. 2.6e+02;  
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTAGACGANG 19

DB 112166 AAGAAAAATCTAGACGANG 112148

RESULT 12 AF012746 2052 bp mRNA linear VRT 24-OCT-2000  
LOCUS AF012746/2052bp 2052 bp mRNA linear VRT 24-OCT-2000  
DEFINITION Dantio perlo olfactory receptor protein 13.1 mRNA, complete cds.  
ACCESSION AF012746.1 GI:2331258  
VERSION AF012746.1 GI:2331258  
KEYWORDS  
SOURCE  
ORGANISM  
Dantio perlo.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;  
Cypriniformes; Cyprinidae; Danio.  
REFERENCE  
1 (bases 1 to 2052)  
Barth, A.L., Justice, N.J. and Ngai, J.  
Asynchronous onset of odorant receptor expression in the developing zebrafish olfactory system  
Neuron 16 (1), 23-34 (1996)

JOURNAL MEDLINE 96158919

REFERENCE PUBMED 8562087

AUTHORS Barth, A.L., Dugas, J.C. and Ngai, J.  
TITLE Noncoordinate expression of odorant receptor genes tightly linked in the zebrafish genome

JOURNAL Neuron 19 (2), 359-369 (1997)

MEDLINE 97436752

REFERENCE PUBMED 9292725

AUTHORS Barth, A.L.  
TITLE Direct Submision

JOURNAL Submitted (07-DEC-1995) Molecular and Cellular Biology, University of California, Berkeley, Km. 265 USA, Berkeley, CA 94720, USA

REFERENCE PUBMED 4 (bases 1 to 2052)

AUTHORS Barth, A.L., Dugas, J.C. and Ngai, J.  
TITLE Direct Submision

JOURNAL Submitted (08-JUL-1997) Molecular and Cellular Biology, University of California, Berkeley, Km. 265 USA, Berkeley, CA 94720, USA

REMARK Sequence updated by submitter  
On Oct 24, 2000 this sequence version replaced gi:1151128.

## FEATURES

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1. 2052  
Location/Qualifiers

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BASE COUNT 638 a 319 c 293 g 802 t  
ORIGIN

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Best Local Similarity 90.9%; Pred. No. 6e+02;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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DB 1472 AAGAAAAATCTAGACAGCAA 1451

RESULT 13 SCYPL252C 2209 bp DNA linear PLN 07-AUG-1997  
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ACCESSION Z73608 U00094  
VERSION Z73608.1 GI:1370516  
KEYWORDS  
SOURCE  
ORGANISM  
Saccharomyces cerevisiae.  
Saccharomyces cerevisiae  
Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;  
Saccharomycetales; Saccharomycetaceae; Saccharomycetes.

REFERENCE 1 (bases 1 to 2209)

AUTHORS Pohl, T.M.  
TITLE Unpublished

JOURNAL 2 (bases 1 to 2209)

AUTHORS MIPS.  
TITLE MIPS.

JOURNAL Direct Submission

Submitted (28-MAY-1996) Data collected by MIPS on behalf of the European yeast chromosome XVI sequencing project. MIPS at the Max-Planck-Institut fuer Biochemie, Am Klopferspitz 18a D-82152 Martinsried, FRG; E-mail: Mewes@mps.emblnet.org

Location/Qualifiers

## FEATURES

source

1. 2209  
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/chromosome="XVI"

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ORIGIN

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Best Local Similarity 90.9%; Pred. No. 6e+02;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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DEFINITION S.cerevisiae chromosome XVI reading frame ORF YPL253C.  
ACCESSION Z73609.000094  
VERSION Z73609.1 GI:1370519  
KEYWORDS  
SOURCE Saccharomyces cerevisiae.  
ORGANISM Saccharomyces cerevisiae.  
Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;  
Saccharomycetales; Saccharomycetaceae; Saccharomyces.

REFERENCE 1 (bases 1 to 2411)  
AUTHORS Pohl,T.M.  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 2411)  
AUTHORS MIPS.  
JOURNAL MIPS.  
TITLE Direct Submission  
JOURNAL Submitted (28-MAY-1996) Data collected by MIPS on behalf of the  
European yeast chromosome XVI sequencing project. MIPS at the  
Max-Planck-Institut fuer Biochemie, Am Klopferspitz 18a D-82152  
Martinsried, FRG; E-mail: Mewes@mips.embl.net.org

FEATURES  
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ORIGIN

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Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 1689 AAGAAAAATCCAAACAGCAA 1688

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LOCUS SC38RCXYI/c  
DEFINITION S.cerevisiae DNA (chromosome XVI; 38 Kb).  
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VERSION Z67751.1 GI:1061234  
KEYWORDS gal4 protein; H(+)-transporting ATPase; HSP82 protein; HSP90  
protein; SRP8 protein; SUI3 protein.  
SOURCE Saccharomyces cerevisiae.  
ORGANISM Saccharomyces cerevisiae.  
Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;  
Saccharomycetales; Saccharomycetaceae; Saccharomyces.

REFERENCE 1 (bases 1 to 37808)  
AUTHORS Pohl,T.M.  
JOURNAL Direct Submission  
REFERENCE 2 (bases 1 to 37808)  
AUTHORS Pohl,T.M.  
JOURNAL Submitted (09-NOV-1995) Thomas M. Pohl, GATC GmbH,  
Fritz-Arnold-Str. 23, Konstanz, 78467, Germany

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complement(18399..19418)

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19902..21701

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AYIRYNLLSISERDLFTLHNNOMKLYTSLPSKTYKKEERLYKMLTYLSDIM  
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IEKLSKKIIPITQIKLNLEPLPKMLPIPSKPTLPDLAFNITTDKQPSASQVKS

Query Match 85.5%: Score 18.8; DB 8; Length 37808;  
Best Local Similarity 90.9%; Pred. No. 3.8e+02;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTAGACAGCA 22  
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DB 4025 AAGAAAAATCTAGACAGCA 4004

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Tue Mar 18 16:16:15 2003

us-09-836-439-4.rge

Page 18

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Job time : 548.495 secs

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GenCore version 5.1.4.p5.4578  
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OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 09:42:12 ; Search time 126.086 Seconds  
(without alignments)  
392.938 Million cell updates/sec

Title: US-09-836-439-4

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Sequence: 1 aagaaaaatctagacaagca 22

Scoring table: IDENTITY\_NUC  
Gapop 10.0, Gapext 1.0

Searched: 2185239 seqs, 112599159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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4	17.4	79.1	412	22	ABAI8136
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8	17.2	78.2	674	22	AAS30251
9	17.2	78.2	674	22	AAS26974

10	17.2	78.2	674	22	AAS26975	Human genomic DNA
11	17.2	78.2	674	22	AAS33485	DNA encoding human
12	17.2	78.2	674	22	AAS33486	DNA encoding human
13	17.2	78.2	782	24	ABD37775	Extended sequence
14	17.2	78.2	786	24	ABQ73702	Human colon specific
15	17.2	78.2	831	24	ABQ43712	Oligonucleotide fo
16	17.2	78.2	831	24	ABQ43713	Oligonucleotide fo
17	17.2	78.2	1372	23	ABV25096	Human prostate exp
18	17.2	78.2	1509	24	ABQ73703	Human colon specif
19	17.2	78.2	1588	21	AAC35087	Arabidopsis thalia
20	17.2	78.2	1622	22	AAK83087	Human immune/haema
21	17.2	78.2	2713	23	ABL24452	Drosophila melanog
22	17.2	78.2	2961	24	ABQ77508	Human cytokine rec
23	17.2	78.2	2963	24	ABQ77508	Human cytokine rec
24	17.2	78.2	3138	23	ABL24782	Drosophila melanog
25	17.2	78.2	3729	22	AAE77688	Human wild-type fc
26	17.2	78.2	4081	23	ABL16178	Drosophila melanog
27	17.2	78.2	4131	22	AAE77689	Human variant Peep
28	17.2	78.2	4258	22	AAE67062	Human immune/haema
29	17.2	78.2	5552	21	AAV58309	Human immune/haema
30	17.2	78.2	6716	24	ABL33783	p1p/pycsp.1 plasm
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32	17.2	78.2	7202	22	ABAI9579	Human immune syst
33	17.2	78.2	9615	22	AAK71750	Human immune/haema
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35	17.2	78.2	11298	19	AAV54661	Human high affinity
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37	17.2	78.2	11298	21	AAA34815	Human adenosine re
38	17.2	78.2	11298	22	AAE92144	Human ICBRI gene S
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#### ALIGNMENTS

RESULT 1  
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XX  
AC ABO69157:  
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DT 29-AUG-2002 (first entry)  
XX  
DE Listeria monocytogenes 4b contig DNA sequence #1923.  
XX  
KW Antibacterial; Listeria; food contamination; mutational analysis;  
XX infection; ds.  
XX  
XX Listeria monocytogenes 4b.  
OS WO200228891-A2.  
XX  
XX  
XX  
XX 11-APR-2002.  
XX  
XX 04-OCT-2001; 2001WO-FR03061.  
XX  
XX 04-OCT-2000; 2000FR-0012697.  
XX  
XX (INSP) INST PASTEUR.  
XX (CNRS) CNRS CENT NAT RECH SCT.  
XX  
XX Kunst F, Glaser P;  
PI WPI; 2002-332479/37.  
XX  
XX New genomic sequences from Listeria species, useful for detection,  
XX treatment and prevention of infection, also related polypeptides,  
PT

PT antibodies and modulators -  
XX  
PS Claim 14; SEQ ID 1970; 180pp; French.  
CC The present invention relates to nucleic acid sequences  
CC (AB067188-AB071212) from *Listeria* sp. The sequences are useful as probes  
CC and primers for identification and/or detection of *Listeria* (e.g. as  
CC contaminants in foods, or mutational analysis) and for analysis of  
CC gene expression. Proteins encoded by the nucleic acid sequences can be  
CC used to screen for compounds that modulate gene expression, replication  
CC and pathogenicity of *Listeria* (potential therapeutic agents), also for  
CC treating infections by *Listeria*, and are useful as immunogens in  
CC anti-*Listeria* vaccines.  
CC Note: The sequence data for this patent did not form part  
CC of the printed specification, but was obtained in electronic format  
CC directly from WIP0 at ftp.wipo.int/pub/published\_pdt\_sequences.  
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KW antiparkinsonian; antischistosomal; antianemic; antidiabetic; cancer;  
KW antineoplastic; hepatotropic; cerebroprotective; antineoplastic;  
KW antiallergic; antidiabetic; antileukemic; anticonvulsant; antifungal;  
KW antiparasitic; cardiac; immune disorder; cardiovascular disorder;  
KW neurological disease; infection; nephrotropic; gene therapy; vaccine; ss.  
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PR 05-JAN-2001; 2001US-0259678.  
  
(HUMA-) HUMAN GENOME SCI INC.  
PA Rosen CA, Barash SC, Ruben SM;  
XX  
XX  
XX WPI: 2001-541565/60.  
DR P-PSDB; ABA16377.  
XX  
XX Nucleic acids encoding 3224 human nervous system antigen polypeptides,  
PT useful for preventing, diagnosing and/or treating nervous system  
PT cancers and metastases -  
XX  
PS Claim 1; SEQ ID NO 1710; 1701tp + Sequence Listing; English.  
XX  
XX The invention relates to novel genes (ABA11004-ABA21534) and proteins  
CC (ABA1678-ABA16001) useful for preventing, treating or ameliorating  
CC medical conditions e.g. by protein or gene therapy. The genes are  
CC isolated from a range of human tissues disclosed in the specification.  
CC The nucleic acids, proteins, antibodies and (ant)agonists are useful  
CC in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast  
CC and ovarian cancer and other cancers of the adrenal gland, bone, bone  
CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;  
CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune  
CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's  
CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative  
CC colitis; (c) cardiovascular disorders such as myocardial ischaemia;  
CC (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and  
CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal  
CC and parasitic infections.

CC Note: The sequence data for this patent did not form part of the  
CC printed specification, but was obtained in electronic format directly  
CC from WIPO at ftp.wipo.int/pub/published\_pcr\_sequences.  
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KW immunosuppressive; antiinflammatory; anti-HIV; antibacterial; vulnerary;  
KW antiparkinsonian; antistickling; antianaemic; antirheumatic; cancer;  
KW antirheumatic; hepatotropic; cerebroprotective; antiinflammatory;  
KW antiallergic; antidiabetic; antitumor; anticonvulsant; antifungal;  
KW antiparasitic; cardiac; immune disorder; cardiovascular disorder;  
KW neurological disease; infection; nephrotropic; gene therapy; vaccine; ds.  
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(HUMA-) HUMAN GENOME SCI INC.

Rosen CA, Barash SC, Ruben SM;

WPI; 2001-541565/60.

PT Nucleic acids encoding 3224 human nervous system antigen polypeptides  
PT useful for preventing, diagnosing and/or treating nervous system  
PT cancers and metastases -

Disclosure; SEQ ID NO 10466; 1701pp + Sequence Listing; English.

CC The invention relates to novel genes (ABAI1004-ABA21534) and proteins  
CC (ABAI4678-ABAI8001) useful for preventing, treating or ameliorating  
CC medical conditions e.g. by protein or gene therapy. The genes are

CC isolated from a range of human tissues disclosed in the specification  
CC The nucleic acids, proteins, antibodies and (ant)agonists are useful

CC in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast  
CC and ovarian cancer and other cancers of the adrenal gland, bone, bone

cc marrow, breast, gastrointestinal tract, liver, lung, or urogenital;  
cc (b) immune disorders e.g. Addison's disease, allergies, autoimmune

CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's  
CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative

cc colitis; (c) cardiovascular disorders such as myocardial ischaemias;  
cc (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and

epilepsy; and (4) infectious diseases such as viral, bacterial, fungal and parasitic infections.

CC Note: The sequence data for this patent did not form part of the  
CC printed specification, but was obtained in electronic format directly  
CC from the patent office.

from WIPO at [http://wipo.int/pub/published\\_pct\\_sequences](http://wipo.int/pub/published_pct_sequences).

Sequence 412 BP; 137 A; 69 C; 71 G; 135 T; 0 other;

Query Match	79.18;	Score 17.4;	DB 22;	Length 412;
Best Local Similarity	94.7%;	Pred. No. 3.8e+02;		

Matches	18;	Conservative	0;	Mismatches	1;	Indels	0;	Gaps	0;
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QY 1 AAGAAAAATCTAGACAAG 19  
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Db 220 AAGAAATATCTAGACAAG 202

RESULT 4  
ABAl8136/C  
ID ABAl8136 standard; DNA: 412 BP.  
XX  
AC ABAl8136;  
XX  
DT 23-JAN-2002 (first entry)  
XX  
DE Human nervous system related polynucleotide SEQ ID NO 10467.  
XX  
KW Human, noctropic; neuroprotective; cytoskeletal; dermatological; virocidic;  
KW immunosuppressive; antihistaminic; anti-HIV; antibacterial; vulnerary;  
KW antiparkinsonian; antiskilling; antianaemic; antidiabetic; cancer;  
KW antirheumatic; hepatotropic; cerebroprotective; antiinflammatory;  
KW antiallergic; antidiabetic; antileucic; anticonvulsant; antifungal;  
KW antiparasitic; cardiac; immune disorder; cardiovascular disorder;  
KW neurological disease; infection; nephrotropic; gene therapy; vaccine; ds.  
XX  
OS Homo sapiens.  
XX  
PN MO200159063-A2.  
XX  
PD 16-AUG-2001.  
XX  
FE 17-JAN-2001; 2001MO-US01334.  
XX  
XX 31-JAN-2000; 2000US-0179065.  
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PR 17-NOV-2000; 2000US-0249264.

[illegible]

XX	OS	Homo sapiens.
XX	PN	WO200160860-A2.
XX	PD	23-AUG-2001.
XX	PF	20-FEB-2001; 2001WO-USO5171.
XX	PR	17-FEB-2000; 2000US-183319P.
XX	PR	16-MAR-2000; 2000US-189862P.
XX	PR	25-MAY-2000; 2000US-207454P.
XX	PR	09-JUN-2000; 2000US-211314P.
XX	PR	18-JUL-2000; 2000US-218007P.
XX	PR	13-DEC-2000; 2000US-255281P.
XX	PA	(MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
XX	P1	Schlegel R, Endege WO, Monahan JE;
XX	P1	WPI; 2001-662795/76.
XX	PT	Novel isolated nucleic acid molecule associated with cancerous state of prostate cells and correlating with presence of prostate cancer, useful for detecting presence of prostate cancer, stage of prostate cancer -
XX	PT	
XX	PS	Claim 1; Page 7745-7746; 1150pp; English.
XX	CC	The invention relates to an isolated nucleic acid molecule (I) comprising
XX	CC	a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the
XX	CC	specification or its complement. (I) is useful for:
XX	CC	(a) assessing whether a patient is afflicted with prostate cancer;
XX	CC	(b) monitoring the progression of prostate cancer in a patient;
XX	CC	(c) assessing the efficacy of a test compound to inhibit prostate
XX	CC	cancer in a patient;
XX	CC	(d) assessing the efficacy of a therapy for inhibiting prostate cancer
XX	CC	in a patient;
XX	CC	(e) selecting a composition for inhibiting prostate cancer in a patient;
XX	CC	(f) assessing the prostate cell carcinogenic potential of a compound;
XX	CC	(g) determining whether prostate cancer has metastasized in a patient;
XX	CC	(h) assessing the aggressiveness or indolence of prostate cancer in a
XX	CC	patient;
XX	CC	(I) is also useful as a pharmacodynamic or pharmacogenomic marker.
XX	CC	
XX	SQ	Sequence 434 BP; 130 A; 78 C; 76 G; 150 T; 0 other;
XX		
XX		Query Match 78.2%; Score 17.2; DB 23; Length 434;
XX		Best Local Similarity 86.4%; Pred.No.4.6e+02;
XX		Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0.
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DB		240 AAGAAAAAAATCTGAAGAAGAAA 219
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XX	AC	ABV58334;
XX	DT	13-SEP-2002 (first entry)
XX		
XX	DE	Human prostate expression marker CDNA 58325.
XX	KX	
XX	KV	Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
XX	KV	pharmacogenomic marker; gene; ss.
XX	OS	Homo sapiens.
XX	XX	
XX	PN	WO200160860-A2.
XX	PD	23-AUG-2001.
XX	XX	

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XX		
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 PR 08-DEC-2000; 2000US-0251989.  
 PR 11-DEC-2000; 2000US-0254097.  
 PR 05-JAN-2001; 2001US-0259678.  
 XX (HUMA-) HUMAN GENOME SCI INC.  
 PA Rosen CA, Barash SC, Ruben SM;  
 XX WPI, 2001-488787/53.  
 XX

PR New polynucleotides and polypeptides, useful for diagnosing, treating,  
 PT preventing or prognosing e.g. kidney, cardiovascular, blood,  
 PT electrolyte imbalance or neoplastic disorders, autoimmune diseases,  
 PT cancers  
 XX  
 PS Claim 1; SEQ ID No 168; 506bp; English.  
 XX  
 CC The invention relates to novel nucleic acids and polypeptides useful for  
 CC diagnosing, treating, preventing and/or prognosing disorders related to  
 CC these polypeptides. The polynucleotides are especially useful in the  
 CC diagnosis, prognosis, prevention and/or treatment of diseases which  
 CC include kidney disorders (e.g. renal failure or nephritis),  
 CC cardiovascular disorders (e.g. hypertension or myocardial infarction),  
 CC blood disorders (e.g. anaemia or blood coagulation disorders),  
 CC electrolyte imbalance disorders (e.g. hyponatraemia or hyperkalaemia),  
 CC neoplastic disorders (e.g. nephroma or renal cell cancer), autoimmune  
 CC diseases, cancers, inflammatory diseases, reproductive system  
 CC disorders, endocrine disorders, neural activity and neurological  
 CC disorders, wound healing and respiratory disorders. AAS30251  
 CC represent the novel human renal and cardiovascular-associated nucleic  
 CC acid sequences of the invention. Note: The sequence data for this  
 CC patent did not form part of the printed specification, but was obtained  
 CC in electronic format directly from WIPO at:  
 CC ftp.wipo.int/pub/published\_pct\_sequences.  
 CC  
 SQ Sequence 674 BP; 144 A; 184 C; 145 G; 201 T; 0 other;  
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 Db 137 AAGAAAAAATCTAGACAGCAA 158  
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 XX  
 AC AAS30251;  
 XX  
 DT 21-NOV-2001 (first entry)  
 XX  
 DE DNA encoding renal and cardiovascular-associated protein, Seq ID 169.  
 XX  
 KW Human; antiinflammatory; neuroprotective; immunomodulator; vulnery;  
 KW cardiovascular; cytosolic; nephrotropic; antinaemic; nephritis;  
 KW immunosuppressive; kidney disorder; renal failure; hypertension;  
 KW cardiovascular disorder; myocardial infarction; blood disorder; anaemia;  
 KW blood coagulation disorder; electrolyte imbalance disorder; cancer;  
 KW hypotatraemia; hyperkalaemia; neoplastic disorder; nephroma;  
 KW autoimmune disease; inflammatory disease; reproductive system disorder;  
 KW endocrine disorder; neural activity; neurological disorder;  
 KW wound healing; respiratory disorder; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200155328-A2.  
 XX  
 PD 02-AUG-2001.  
 XX  
 PE 17-JAN-2001; 2001MO-US01359.  
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 PR 07-JUN-2000; 2000US-0209467.



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XX	05-JAN-2001;	2001US-0239678.
PA	(HUMA-) HUMAN GENOME SCI INC	
PI	Rosen CA, Barash SC, Ruben	
XX	WPI; 2001-488787/53.	
XX	New polynucleotides and poly-	
PT	preventing or prognosing e.g	
PT	electrolyte imbalance or neo-	
XX	cancers -	
PS	Claim 1; SEQ ID No 169; 506pp)	
CC	The invention relates to novel	
CC	diagnosing, treating, preven	
CC	these polypeptides. The poly	
CC	diagnosis, prognosis, preven	
CC	include kidney disorders (e.i.	

CC cardiovascular disorders (e.g. hypertension or myocardial infarction),  
CC blood disorders (e.g. anaemia or blood coagulation disorders),  
CC electrolyte imbalance disorders (e.g. hyponatraemia or hyperkalaemia),  
CC neoplastic disorders (e.g. nephroma or renal cell cancer), autoimmune  
CC diseases, cancers, inflammatory diseases, reproductive system  
CC disorders, endocrine disorders, neural activity and neurological  
CC disorders, wound healing and respiratory disorders. AAS30165-AAS30251  
CC represent the novel human renal and cardiovascular-associated nucleic  
CC acid sequences of the invention. Note: The sequence data for this  
CC patent did not form part of the printed specification, but was obtained  
CC in electronic format directly from WIPD at:  
CC ftp.wipo.int/pub/published\_pct\_sequences.  
CC  
CC  
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KW Human; immunosuppressive; antiarthritic; ds; antirheumatic;  
KW cytoskeletal; cardiac; vasotropic; cerebroprotective; nootropic;  
KW neuroprotective; antibiotic; virucide; fungicide; ophthalmological;  
KW vulnerrary; secreted protein; rheumatoid arthritis;  
KW hyperproliferative disorder; cardiovascular disorder; cardiac arrest;  
KW cerebrovascular disorder; cerebral ischemia; angiogenesis;  
KW nervous system disorder; Alzheimer's disease; infection; ocular disorder;  
KW corneal infection; wound healing; epithelial cell proliferation;  
KW skin ageing; food additive; preservative; antiproliferative.  
OS Homo sapiens.  
XX  
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PN WO20015441-A2.  
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 PR 11-DEC-2000; 2000US-0254097.  
 PR 05-JAN-2001; 2001US-0259678.  
 XX (HUMA-) HUMAN GENOME SCI INC.  
 PA Rosen CA, Barash SC, Ruben SM;  
 PI WPI: 2001-476222/51.  
 XX  
 DR Novel polypeptides and polynucleotides useful as diagnostic reagents to  
 XX diagnose diseases or disorders associated with aberrant expression or  
 PT activity of polypeptides, for treating blood clotting disorder,  
 PT haemophilia  
 PS Disclosure: SEQ ID No 310; 601pp; English.  
 XX  
 CC The invention relates to isolated nucleic acid molecules and their  
 CC encoded secreted proteins. The nucleic acids and proteins are used to  
 CC prevent, treat or ameliorate a medical condition in e.g. humans, mice,  
 CC rabbits, goats, horses, cats, dogs, chickens or sheep. They  
 CC are also used in diagnosing a pathological condition or susceptibility  
 CC to a pathological condition. Antibodies to the proteins can also  
 CC be used in alleviating symptoms associated with the disorders and in  
 CC diagnostic immunoassays e.g. radioimmunoassays or enzyme linked  
 CC immunosorbent assays (ELISA). Disorders which are diagnosed or treated  
 CC include autoimmune diseases e.g. rheumatoid arthritis,  
 CC hyperproliferative disorders e.g. neoplasms of the breast or liver,  
 CC cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders  
 CC e.g. cerebral ischaemia, angiogenesis, nervous system disorders e.g.  
 CC Alzheimer's disease, infections caused by bacteria, viruses and fungi  
 CC and ocular disorders e.g. corneal infection, and many other  
 CC disorders listed in the specification. The polypeptides can also  
 CC be used to aid wound healing and epithelial cell proliferation, to

CC prevent skin aging due to sunburn, to maintain organs before  
 CC transplantation, for supporting cell culture of primary tissues, to  
 CC regenerate tissues and in chemotaxis. The polypeptides can also be used  
 CC as a food additive or preservative to increase or decrease storage  
 CC capabilities, fat content, lipid, protein, carbohydrate, vitamins,  
 CC minerals, cofactors and other nutritional components. The present  
 CC sequence is a genomic DNA encoding a partial novel secreted protein of

Query Match 78.2%; Score 17.2; DB 22; Length 674;  
 Best Local Similarity 86.4%; Pred. No. 4.6e+02;  
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Gy 1 AAGAAAAATCTAGACAGCAA 22  
 Db 137 AAGAAAAATTTTAAACAAACA 158

RESULT 10  
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 XX Human; immunosuppressive; antiarthritic; ds; antirheumatic;  
 KW cytototoxic; cardiant; vasotropic; cerebroprotective; nootropic;  
 KW neuroprotective; antibacterial; virucide; fungicide; optalmalogical;  
 KW vulnery; secreted protein; rheumatoid arthritis;  
 KW hyperproliferative disorder; cardiovascular disorder; cardiac arrest;  
 KW cerebrovascular disorder; cerebral ischaemia; angiogenesis;  
 KW nervous system disorder; Alzheimer's disease; infection; ocular disorder;  
 KW corneal infection; wound healing; epithelial cell proliferation;  
 KW skin ageing; food additive; preservative; antiproliferative.  
 XX  
 OS Homo sapiens.  
 XX  
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(HUMA-) HUMAN GENOME SCI INC.

Rosen CA, Barash SC, Ruben SM;

WPI: 2001-476222/51.

Novel polypeptides and polynucleotides useful as diagnostic reagents to diagnose diseases or disorders associated with aberrant expression or activity of polypeptides, for treating blood clotting disorder, haemophilia

Disclosure; SEQ ID No 311; 601pp; English.

The invention relates to isolated nucleic acid molecules and their encoded secreted proteins. The nucleic acids and proteins are used to prevent, treat or ameliorate a medical condition in e.g. humans, mice, rabbits, goats, horses, cats, dogs, chickens or sheep. They are also used in diagnosing a pathological condition or susceptibility to a pathological condition. Antibodies to the proteins can also be used in alleviating symptoms associated with the disorders and in diagnostic immunoassays e.g. radioimmunoassays or enzyme linked immunoassay assays (ELISA). Disorders which are diagnosed or treated include autoimmune diseases e.g. rheumatoid arthritis, hyperproliferative disorders e.g. neoplasms of the breast or liver, cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders e.g. cerebral ischemia, angiogenesis, nervous system disorders e.g. Alzheimer's disease, infections caused by bacteria, viruses and fungi and ocular disorders e.g. corneal infection, and many other disorders listed in the specification. The polypeptides can also be used to aid wound healing and epithelial cell proliferation, to prevent skin aging due to sunburn, to maintain organs before transplantation, for supporting cell culture of primary tissues, to regenerate tissues and in chemotaxis. The polypeptides can also be used as a food additive or preservative to increase or decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamins, minerals, cofactors and other nutritional components. The present sequence is a genomic DNA encoding a partial novel secreted protein of

Query Match 78.2%; Score 17.2; DB 22; Length 674;  
Best Local Similarity 86.4%; Pred. No. 4.6e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 AAGAAAAAATCTAGACAGCAA 22  
|||||  
DB 137 AAGAAAAAATTTAAACAAACAA 158

## RESULT 11

AAS33485

ID AAS33485 standard; DNA; 674 BP.

AC AAS33485;

DT 04-DEC-2001 (first entry)

DE DNA encoding human secreted protein, Seq ID No 768.

XX Immunomodulatory; human immunodeficiency virus; HIV; anaemia; angina;  
XX rheumatoid arthritis; antiarteriosclerotic; cardiant; vascular;  
XX cerebroprotective; thrombolytic; antimicrobial; ophthalmological;  
XX cytoskeletal; Alzheimer's disease; Parkinson's disease; human; cancer;  
XX multiple sclerosis; cancer; hyperproliferative disorder; infection;  
XX Gaucher's disease; neurological disease; cerebrovascular disorder;  
XX thrombosis; wound healing; ds.

OS Homo sapiens.

PN WO200153326-A2.

PD 02-AUG-2001.

PF 17-JAN-2001; 2001WO-US01347.

PR 31-JAN-2000; 2000US-0179065.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Rosen CA, Barash SC, Ruben SM;

DR WPI: 2001-451931/48.

PT New nucleic acids and polypeptides, useful for diagnosing, preventing  
or treating medical conditions -

PS Disclosure: SEQ ID No 768; 753bp; English.

XX The invention relates to novel isolated nucleic acid molecules (I)  
XX encoding human secreted proteins (II). (I) and (II) are used to prevent,  
XX treat or ameliorate a medical condition in e.g. humans, mice, rabbits,  
XX goats, horses, cats, dogs, chickens or sheep. (I) and (II) may be used in  
XX the prevention, treatment and diagnosis of diseases associated with  
XX inappropriate expression of secreted proteins. (I) and complementary  
XX sequences may also be used as DNA probes in diagnostic assays (e.g.  
XX polymerase chain reactions (PCR)) to detect and quantitate the presence  
XX of similar nucleic acid sequences in samples, and so which patients may  
XX be in need of restorative therapy. (II) may also be used as antigens in  
XX the production of antibodies and in assays to identify modulators  
XX (agonists and antagonists) of the expression and activity of the secreted  
XX proteins. The anti-(II) antibodies and antagonists may also be used to  
XX down regulate expression and activity of (II). The anti-(II) antibodies  
XX may also be used as diagnostic agents for detecting the presence of (II)  
XX in samples (e.g. by enzyme linked immunosorbent assay (ELISA)). The  
XX disorders include for example: immune/autoimmune diseases (e.g. HIV  
XX (human immunodeficiency virus) infections, anaemia, rheumatoid arthritis  
XX and multiple sclerosis), cancers and hyperproliferative disorders (e.g.  
XX melanomas, neoplasms of the breast or liver, Sezary syndrome and  
XX Gaucher's disease), neurological diseases (e.g. Alzheimer's disease,  
XX Parkinson's disease and Charcot-Marie-Tooth disease), cardio-/  
XX cerebrovascular disorders (e.g. cardiac arrest, tachycardia,  
XX angina and thrombosis), infections caused by bacteria, viruses and

CC fungi and ocular disorders (e.g. corneal infections). (I) and (II),  
CC agonists, antagonists and antibodies can also be used to promote wound  
CC healing, maintain organs before transplantation, and support cell culture  
CC of primary tissues. AAS33043-AAS33486 represent human secreted protein  
CC coding sequences, PCR primers, and related sequences of the invention.  
CC Note: The sequence data for this patent did not appear in the printed  
CC specification but was obtained in electronic format directly from WIPO  
CC at: ftp.wipo.int/pub/published\_pcr\_sequences.

XX Sequence 674 BP; 144 A; 184 C; 145 G; 201 T; 0 other;

Query Match 78.2%; Score 17.2; DB 22; Length 674;  
Best Local Similarity 86.4%; Pred. No. 4.6e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 AAGAAAAAATCTAGACAGCAA 22  
|||||  
DB 137 AAGAAAAAATTTAAACAAACAA 158

## RESULT 12

AAS33486

ID AAS33486 standard; DNA; 674 BP.

AC AAS33486;

DT 04-DEC-2001 (first entry)

DE DNA encoding human secreted protein, Seq ID No 769.

XX Immunomodulatory; human immunodeficiency virus; HIV; anaemia; angina;  
XX rheumatoid arthritis; antiarteriosclerotic; cardiant; vascular;  
XX cerebroprotective; thrombolytic; antimicrobial; ophthalmological;  
XX cytoskeletal; Alzheimer's disease; Parkinson's disease; human; cancer;  
XX multiple sclerosis; cancer; hyperproliferative disorder; infection;  
XX Gaucher's disease; neurological disease; cerebrovascular disorder;  
XX thrombosis; wound healing; ds.

OS Homo sapiens.

PN WO200153326-A2.

PD 02-AUG-2001.

PF 17-JAN-2001; 2001WO-US01347.

PR 31-JAN-2000; 2000US-0179065.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Rosen CA, Barash SC, Ruben SM;

DR WPI: 2001-451931/48.

PT New nucleic acids and polypeptides, useful for diagnosing, preventing  
or treating medical conditions -

PS Disclosure: SEQ ID No 769; 753bp; English.

XX The invention relates to novel isolated nucleic acid molecules (I)  
XX encoding human secreted proteins (II). (I) and (II) are used to prevent,  
XX treat or ameliorate a medical condition in e.g. humans, mice, rabbits,  
XX goats, horses, cats, dogs, chickens or sheep. (I) and (II) may be used in  
XX the prevention, treatment and diagnosis of diseases associated with  
XX inappropriate expression of secreted proteins. (I) and complementary  
XX sequences may also be used as DNA probes in diagnostic assays (e.g.  
XX polymerase chain reactions (PCR)) to detect and quantitate the presence  
XX of similar nucleic acid sequences in samples, and so which patients may  
XX be in need of restorative therapy. (II) may also be used as antigens in  
XX the production of antibodies and in assays to identify modulators  
XX (agonists and antagonists) of the expression and activity of the secreted  
XX proteins. The anti-(II) antibodies and antagonists may also be used to  
XX down regulate expression and activity of (II). The anti-(II) antibodies

CC may also be used as diagnostic agents for detecting the presence of (II)  
CC in samples (e.g. by enzyme linked immunosorbent assay (ELISA)). The  
CC disorders include for example: immune/autoimmune diseases (e.g. HIV  
CC (human immunodeficiency virus) infections, anemia, rheumatoid arthritis  
CC and multiple sclerosis), cancers and hyperproliferative disorders (e.g.  
CC melanomas, neoplasms of the breast or liver, Sezary syndrome and  
CC Parkinson's disease), neurological diseases (e.g. Alzheimer's disease,  
CC Charcot-Marie-Tooth disease), cardio-  
CC cerebrovascular disorders (e.g. cardiac arrest, tachycardia,  
CC angina and thrombosis), infections caused by bacteria, viruses and  
CC fungi and ocular disorders (e.g. corneal infections). (I) and (II),  
CC antagonists and antibodies can also be used to promote wound  
CC healing, maintain organs before transplantation, and support cell culture  
CC of primary tissues. AAS3043-AAS3486 represent human secreted protein  
CC coding sequences, PCR primers, and related sequences of the invention.  
CC Note: The sequence data for this patent did not appear in the printed  
CC specification but was obtained in electronic format directly from WIPO  
CC at: [ftp.wipo.int/pub/published\\_pct\\_sequences](http://wipo.int/pub/published_pct_sequences).

SO Sequence 674 BP; 144 A; 184 C; 145 G; 201 T; 0 other;

Query Match 78.2%; Score 17.2; DB 22; Length 674;  
Best Local Similarity 86.4%; Pred. No. 4.6e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 AAGAAAAATCTAGACAGCAA 22  
||| ||||| ||| ||| |||  
DB 137 AAGAAAAATTTAAACAAACAA 158

## RESULT 13

AAD37775/c  
ID AAD37775 standard; DNA; 782 BP.

AC AAD37775;

DT 27-AUG-2002 (first entry)

XX Extended sequence for mouse IMX5\_09.

XX Inflammatory bowel disease; IBD; autoimmune disorder; arthritis; allergy;  
KM haematopoietic cell; thrombolytic; blood coagulation disorder; nephritis;  
KM asthma; organ rejection; graft-versus-host disease; inflammation; shock;  
KM nerve disease; Alzheimer's disease; Parkinson's disease; antibacterial;  
KM Huntington's disease; immunosuppressive; sepsis; nephrotropic; nootropic;  
KM neuroprotective; anticonvulsant; gene therapy; mouse; ds.

XX Mus musculus.

XX WO200231116-A2.

XX 18-APR-2002.

XX 11-OCT-2001; 2001WO-US32176.

XX 11-OCT-2000; 2000US-239712P.

XX (DIGI-) DIGITAL GENE TECHNOLOGIES INC.

XX Vlhay JL, Slims JE, Dubose RF, Baum PR, Hasel KW, Hilbush BS;

XX WPI; 2002-426280/45.

XX New polynucleotide associated with inflammatory bowel disease for  
PT treating disorders of the immune system, nervous system, hematopoietic  
PT cells and to modulate inflammation -

XX Claim 1; Page 202-203; 214pp; English.

XX The invention relates to an isolated polynucleotide associated with  
CC inflammatory bowel disease (IBD). The invention is useful for  
CC manufacturing a medicament for use in preventing, treating, modulating,  
CC or ameliorating a medical condition which is IBD. The polypeptide and

CC polynucleotide are useful for treating disorders of the immune system  
CC e.g. autoimmune disorders, deficiencies or disorders of haematopoietic  
CC cells, to modulate hemostatic, or thrombolytic activity, treat blood  
CC coagulation disorders, allergic reactions and conditions, such as asthma,  
CC treat and/or prevent organ rejection or graft-versus-host disease and  
CC modulate inflammation, including inflammation associated with infection,  
CC shock, sepsis, arthritis and nephritis. The invention is useful to  
CC differentiate, proliferate and attract cells, leading to the regeneration  
CC of tissues and to treat central and peripheral nerve diseases e.g.  
CC Alzheimer's disease, Parkinson's disease, and Huntington's disease. The  
CC invention is useful in gene therapy. The present sequence is an extended  
CC sequence for mouse IMX5\_09 which is used in the exemplification of the  
CC invention.

SO Sequence 782 BP; 212 A; 180 C; 138 G; 252 T; 0 other;

Query Match 78.2%; Score 17.2; DB 24; Length 782;  
Best Local Similarity 86.4%; Pred. No. 4.7e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 AAGAAAAATCTAGACAGCAA 22  
||| ||||| ||| ||| |||  
DB 393 AAGAAAAATCTAAACAAACAA 372

## RESULT 14

ABO73702/c  
ID ABO73702 standard; cDNA; 786 BP.

AC ABO73702;

DT 07-OCT-2002 (first entry)

XX Human colon specific nucleic acid (CSNA) SEQ ID NO:8.

XX Human; colon specific nucleic acid; colon specific polypeptide; CSP;  
KM CSNA; colon specific gene; CSG; colon cancer; gene therapy; vaccine;  
KM cytosolic; gene; ss.

XX Homo sapiens.

XX WO200248370-A2.

XX 20-JUN-2002.

XX 30-OCT-2001; 2001WO-US51341.

XX 31-OCT-2000; 2000US-244717P.

XX (DIAD-) DIADEXUS INC.

XX Sun Y, Reclon H, Ghosh MG, Liu C;

XX WPI; 2002-583520/62.

XX Colon specific polypeptides and polynucleotides useful for detecting,  
PT diagnosing, monitoring, treating, staging and predicting cancers in  
PT humans having cancer and non-cancerous colon disease -

XX Claim 1; Page 153; 243pp; English.

XX ABO73695 to ABO73841 represent human colon specific nucleic acid (CSNA)  
CC sequences, and ABO73826 to ABO73928 represent human colon specific  
CC polypeptide (CSP) sequences from the present invention. CSNA and CSP  
CC sequences have cytostatic activity, and can be used in gene therapy,  
CC antisense therapy and in vaccines. CSNA and CSP sequences can be used  
CC for diagnosing and monitoring the presence and metastases of colon  
CC cancer in a patient, by determining an amount of CSP or CSNA in a sample  
CC of a patient, and comparing it to the amount of colon specific marker  
CC in a normal control, where a difference in the amount of the nucleic  
CC acid or the polypeptide in the sample compared to that of normal control  
CC is associated with presence of colon cancer. CSP and CSNA sequences can  
CC be used for producing engineered colon tissue for treatment and research.

CC CSNA sequences are useful for producing transgenic animals and cells  
 CC and also in gene therapy.

XX  
 SQ Sequence 786 BP; 256 A; 118 C; 140 G; 272 T; 0 other;

Query Match 78.2%; Score 17.2; DB 24; Length 786;  
 Best Local Similarity 86.4%; Pred. No. 4.7e+02;

Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTAGACACGCA 22  
 ||| ||||| ||||| |||||  
 Db 715 ATGAGAAAAATCTAACAACGCA 694

RESULT 15  
 ABQ43712/C  
 ID ABQ43712 standard; DNA; 831 BP.

AC ABQ43712;

DT 12-JUL-2002 (first entry)

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 30303.

XX  
 KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;  
 KW drug; side effect; cancer; central nervous system; cardiovascular;  
 KW gastrointestinal; respiratory system; single nucleotide polymorphism;  
 KW SNP; cell differentiation; ds.

OS Homo sapiens.

XX MO200218632-A2.

PD 07-MAR-2002.

PF 01-SEP-2001; 2001WO-EP10074.

PR 01-SEP-2000; 2000DE-1043826.  
 PR 05-SEP-2000; 2000DE-1044543.

PA (EPIC-) EPICENOMICS AG.

PI Olek A, Piepenbrock C, Berlin K, Guetig D;

DR WPI; 2002-371829/40.

PT Determining the degree of cytosine methylation in genomic DNA, useful  
 PT for diagnosis and prognosis, comprises selective hybridization of  
 PT amplicons from chemically treated DNA -

PS Claim 12: 56pp + Sequence Listing; 56pp; German.

XX  
 CC This invention describes a novel method for determining the degree of  
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a  
 CC genomic sample of DNA. The sample is treated chemically to convert in a  
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic  
 CC DNA that contains the target C is amplified to form a labeled amplicon.  
 CC The amplicon is hybridized to two classes, each with at least one  
 CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers  
 CC and the degree of hybridization to both classes is determined from the  
 CC label on the amplicon. From the ratio of labels hybridized to the two  
 CC classes of oligomers, the degree of methylation is calculated. The method  
 CC is used: (i) for diagnosis and/or prognosis of side effects of  
 CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders  
 CC of the central nervous, cardiovascular, gastrointestinal and respiratory  
 CC systems etc., particularly by detecting mutations or single nucleotide  
 CC polymorphisms (SNP's); and (ii) for differentiation of cell or tissue  
 CC types and for investigating cell differentiation. The method allows the  
 CC methylation status of many C residues to be determined simultaneously.  
 CC ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the  
 CC method for determining the degree of cytosine methylation described in  
 CC the disclosure of the invention.

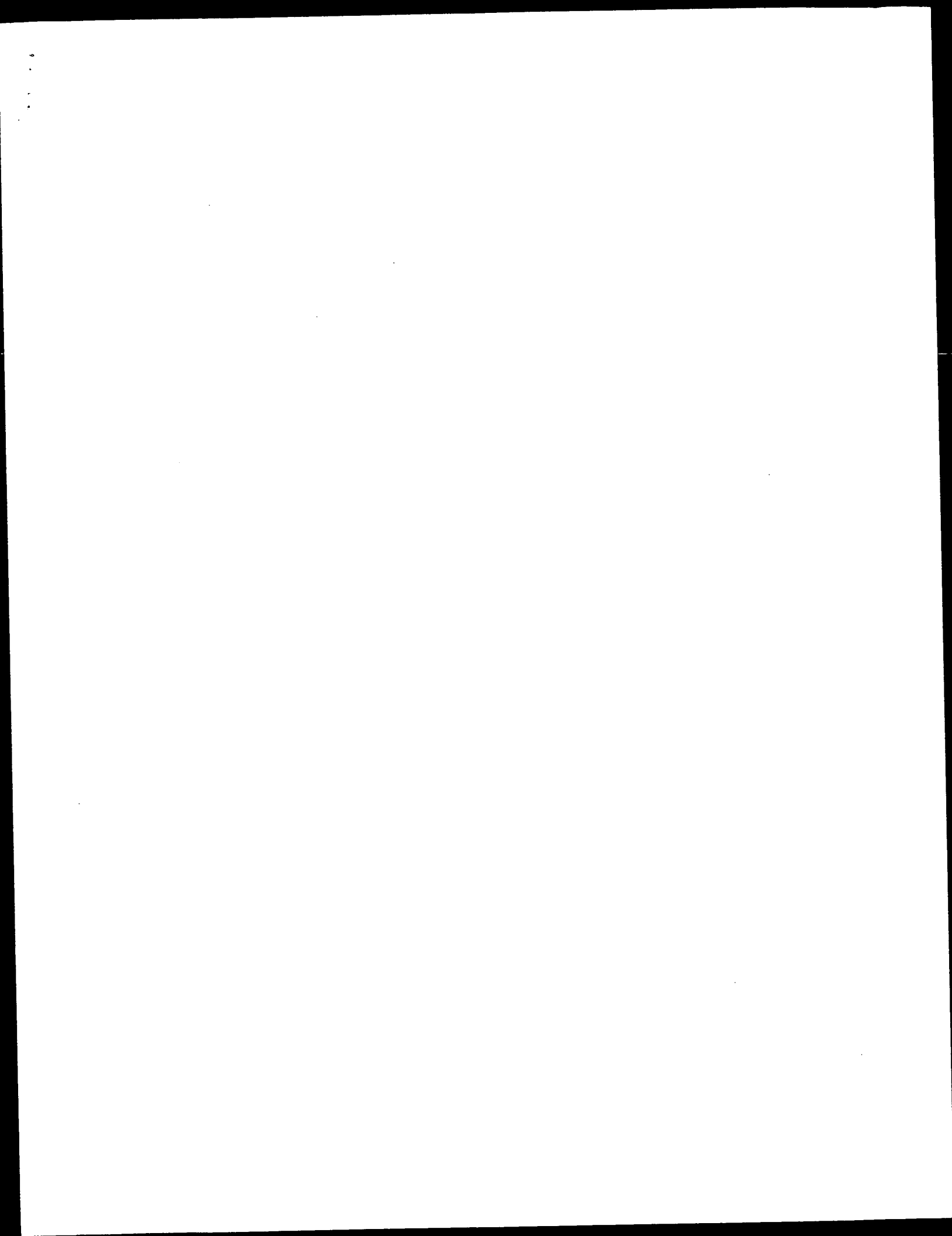
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Query Match 78.2%; Score 17.2; DB 24; Length 831;  
 Best Local Similarity 86.4%; Pred. No. 4.7e+02;

Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTAGACACGCA 22  
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 Db 771 AAAAAAAATCTAACAACGCA 750

Search completed: March 17, 2003, 10:50:47  
 Job time: 128.253 secs





GenCore version 5.1.4-p5-4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 10:23:12 ; Search time 777.688 Seconds

(without alignments)  
458.154 Million cell updates/sec

Title: US-09-836-439-4

Perfect score: 22  
Sequence: 1 aagaaaaatcagaacaa 22

Scoring table: IDENTITY NUC  
Gapop 10.0, Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

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6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hc:\*  
9: gb\_estl:\*  
10: gb\_est2:\*  
11: gb\_hc:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
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16: em\_estom:\*  
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18: em\_gss\_hum:\*  
19: em\_gss\_low:\*  
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21: em\_gss\_vrt:\*  
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26: em\_gss\_pro:\*  
27: em\_gss\_rtd:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20.4	92.7	342	10	AM740409
2	20.4	92.7	342	12	BS360474
3	20.4	92.7	360	10	AM740410
4	18.8	85.5	224	10	BB016617
5	18.8	85.5	424	17	BB670801
6	18.8	85.5	449	17	AZ009207

C	7	18.8	85.5	476	10	BB768795	BB768795
C	8	18.8	85.5	505	12	BF004802	BF004802
C	9	18.8	85.5	528	10	AA447445	AA447445
C	10	18.8	85.5	544	17	AA014714	AA014714
C	11	18.8	85.5	549	17	BB282218	BB282218
C	12	18.8	85.5	567	17	AA0534621	AA0534621
C	13	18.8	85.5	568	17	AA2506877	AA2506877
C	14	18.8	85.5	597	17	AA0534617	AA0534617
C	15	18.8	85.5	710	17	BB552202	BB552202
C	16	18.8	85.5	738	17	AA2572374	AA2572374
C	17	18.8	85.5	767	17	CNS07C7K	CNS07C7K
C	18	18.8	85.5	1059	17	CNS06H8C	CNS06H8C
C	19	18.8	85.5	1330	12	BF164624	BF164624
C	20	18.4	83.6	479	17	CNS010U1	CNS010U1
C	21	18.8	81.8	507	10	AA828026	AA828026
C	22	18.8	81.8	582	10	AA871607	AA871607
C	23	18.8	81.8	597	10	AA870709	AA870709
C	24	18.8	81.8	624	10	AA735590	AA735590
C	25	18.8	81.8	651	10	AA735590	AA735590
C	26	17.8	80.9	229	10	AA380205	AA380205
C	27	17.8	80.9	280	10	BB381647	BB381647
C	28	17.8	80.9	283	10	BB106733	BB106733
C	29	17.8	80.9	318	9	AL837465	AL837465
C	30	17.8	80.9	318	13	BI034283	BI034283
C	31	17.8	80.9	501	10	AA873095	AA873095
C	32	17.8	80.9	510	17	AZ035704	AZ035704
C	33	17.8	80.9	527	10	AA873962	AA873962
C	34	17.8	80.9	533	14	BO234625	BO234625
C	35	17.8	80.9	539	17	BA427923	BA427923
C	36	17.8	80.9	558	10	AA034365	AA034365
C	37	17.8	80.9	632	17	AZ417128	AZ417128
C	38	17.8	80.9	638	10	AA872452	AA872452
C	39	17.8	80.9	656	17	BB059762	BB059762
C	40	17.8	80.9	666	10	BB641601	BB641601
C	41	17.8	80.9	666	17	BB717507	BB717507
C	42	17.8	80.9	670	17	DR10N24S	DR10N24S
C	43	17.8	80.9	703	10	AA868791	AA868791
C	44	17.8	80.9	775	17	BB705808	BB705808
C	45	17.8	80.9	781	17	BB059760	BB059760

## ALIGNMENTS

RESULT 1  
LOCUS AM740409 342 bp mRNA linear EST 27-APR-2000  
DEFINITION BR110552 Blomphalaria glabrata (BS-90)-unexposed lambda zap library  
ACCESSION Blomphalaria glabrata cDNA clone RBG1155TR, mRNA sequence.  
VERSION AM740409.1 GI:7651688  
KEYWORDS EST.

## SOURCE

ORGANISM Blomphalaria glabrata

REFERENCE 1 (bases 1 to 342)  
Eukaryote: Metazoa: Mollusca: Gastropoda: Pulmonata; Basommatophora  
Planorbidae: Blomphalaria.

## AUTHORS

Raghavan, N., Miller, A., Gardner, M., Kerlavage, A.R., Fitzgerald, P.C.,  
Lewis, F.A., and Knight, M.

## TITLE

Genes expressed by the hemocytes of Blomphalaria glabrata before  
and after exposure to miracidia

## JOURNAL

## COMMENT

Unpublished (2000)  
Contact: Raghavan N  
Biomedical Research Institute  
1211 Parklawn Dr., Rockville, MD 20852, USA  
Tel: 301-881-3300 ext.128  
Fax: 301-770-4756  
Email: nkr@helix.nih.gov, snallstrule@aol.com.

## FEATURES

source  
1..342  
Location/Qualifiers  
/organism="Blomphalaria glabrata"  
/strain="BS-90"  
/db\_xref="taxon:6526"

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/clone="RBGIH55TR"
/clone.lib="Biomphalaria glabrata (BS-90) -unexposed Lambda
zap library"
/sex="hermaphrodite"
/cell_type="Hemocyte"
/lab_host="Laboratory host"
/note="Vector: pBluescript SK-; Site_1: EcoRI; Site_2:
XhoI; Total RNA was isolated from the hemocytes of
unexposed Biomphalaria glabrata (BS-90) snails and first
strand cDNA synthesized using an oligo-dT primer-linker
(XhoI). Second strand synthesis was followed by the
ligation of EcoRI adaptors. Following digestion with XhoI,
the completed, directional cDNA was cloned into Uni-ZAP
XR phagemid vector by Stratagene.
BASE COUNT      132 a      68 c      62 g      80 t
ORIGIN
Query Match      92.7%; Score 20.4; DB 10; Length 342;
Best Local Similarity 95.5%; Pred. No. 1.1e+03;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY      1 AAGAAAAATCTAGACAGCAA 22
|||||
Db      174 AAGAAAAATCTAGACAGCAA 195

RESULT 2
BG360474      342 bp      mRNA      linear      EST 07-MAR-2001
LOCUS      BRL10646 Biomphalaria glabrata (BS-90) -unexposed Lambda zap library
DEFINITION      Biomphalaria glabrata cDNA clone RBGIU88TR, mRNA sequence.
ACCESSION      BG360474
VERSION      BG360474.1 GI:13243488
KEYWORDS      EST.
SOURCE      bloodfluke planorb.
ORGANISM      Biomphalaria glabrata
Eukaryota; Metazoa; Mollusca; Gastropoda; Pulmonata; Basommatophora
; Planorbidae; Biomphalaria.
REFERENCE      1 (bases 1 to 342)
AUTHORS      Raghavan,N., Miller,A., Gardner,M., Kerlavage,A.R., Fitzgerald,P.C.,
Lewis,F.A. and Knight,M.
Genes expressed by the hemocytes of Biomphalaria glabrata before
and after exposure to miracidia
Unpublished (2000)
JOURNAL      Contact: Raghavan N
Biomedical Research Institute
12111 Parklawn Dr., Rockville, MD 20852, USA
Tel: 301-881-3300 ext.128
Fax: 301-770-4756
Email: nkr@helix.nih.gov, snailstrule@aol.com.
COMMENT
FEATURES
source
1. .342
Location/Qualifiers
/organism="Biomphalaria glabrata"
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/db_xref="taxon:6526"
/clone="RBGI88TR"
/clone.lib="Biomphalaria glabrata (BS-90) -unexposed Lambda
zap library"
/sex="hermaphrodite"
/cell_type="Hemocyte"
/lab_host="Laboratory host"
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XhoI; Total RNA was isolated from the hemocytes of
unexposed Biomphalaria glabrata (BS-90) snails and first
strand cDNA synthesized using an oligo-dT primer-linker
(XhoI). Second strand synthesis was followed by the
ligation of EcoRI adaptors. Following digestion with XhoI,
the completed, directional cDNA was cloned into Uni-ZAP
XR phagemid vector by Stratagene.
BASE COUNT      133 a      68 c      63 g      78 t
ORIGIN
Query Match      92.7%; Score 20.4; DB 12; Length 342;

```

```

Best Local Similarity 95.5%; Pred. No. 1.1e+03;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY      1 AAGAAAAATCTAGACAGCAA 22
|||||
Db      174 AAGAAAAATCTAGACAGCAA 195

RESULT 3
AM740410      360 bp      mRNA      linear      EST 27-APR-2000
LOCUS      BRL10553 Biomphalaria glabrata (BS-90) -unexposed Lambda zap library
DEFINITION      Biomphalaria glabrata cDNA clone RBGIH56TR, mRNA sequence.
ACCESSION      AM740410
VERSION      AM740410.1 GI:7651689
KEYWORDS      EST.
SOURCE      bloodfluke planorb.
ORGANISM      Biomphalaria glabrata
Eukaryota; Metazoa; Mollusca; Gastropoda; Pulmonata; Basommatophora
; Planorbidae; Biomphalaria.
REFERENCE      1 (bases 1 to 360)
AUTHORS      Raghavan,N., Miller,A., Gardner,M., Kerlavage,A.R., Fitzgerald,P.C.,
Lewis,F.A. and Knight,M.
Genes expressed by the hemocytes of Biomphalaria glabrata before
and after exposure to miracidia
Unpublished (2000)
JOURNAL      Contact: Raghavan N
Biomedical Research Institute
12111 Parklawn Dr., Rockville, MD 20852, USA
Tel: 301-881-3300 ext.128
Fax: 301-770-4756
Email: nkr@helix.nih.gov, snailstrule@aol.com.
COMMENT
FEATURES
source
1. .360
Location/Qualifiers
/organism="Biomphalaria glabrata"
/strain="BS-90"
/db_xref="taxon:6526"
/clone="RBGIH56TR"
/clone.lib="Biomphalaria glabrata (BS-90) -unexposed Lambda
zap library"
/sex="hermaphrodite"
/cell_type="Hemocyte"
/lab_host="Laboratory host"
/note="Vector: pBluescript SK-; Site_1: EcoRI; Site_2:
XhoI; Total RNA was isolated from the hemocytes of
unexposed Biomphalaria glabrata (BS-90) snails and first
strand cDNA synthesized using an oligo-dT primer-linker
(XhoI). Second strand synthesis was followed by the
ligation of EcoRI adaptors. Following digestion with XhoI,
the completed, directional cDNA was cloned into Uni-ZAP
XR phagemid vector by Stratagene.
BASE COUNT      137 a      72 c      62 g      86 t      3 others
ORIGIN
Query Match      92.7%; Score 20.4; DB 10; Length 360;
Best Local Similarity 95.5%; Pred. No. 1.1e+03;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY      1 AAGAAAAATCTAGACAGCAA 22
|||||
Db      174 AAGAAAAATCTAGACAGCAA 195

RESULT 4
BB016617      224 bp      mRNA      linear      EST 22-JUN-2000
LOCUS      BB016617 RIKEN full-length enriched, adult male testis (DH10B) Mus
DEFINITION      musculus cDNA clone 4930563D02 3', mRNA sequence.
ACCESSION      BB016617
VERSION      BB016617.1 GI:8187765
KEYWORDS      EST.
SOURCE      house mouse.
ORGANISM      Mus musculus

```

Eukaryota, Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sclurognathi; Muridae; Murinae; Mus 1 (bases 1 to 224)  
Konno, H., Aizawa, K., Akahira, S., Akiyama, T., Arahata, T. *Cranioc*

Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi: Mammalia: Eutheria: Rodentia: Scurionathia: Muridae; Murine; Mus. 1 (bases 1 to 224)

Konno, H., Aizawa, K., Akahira, S., Akiyama, U., Arakawa, T., Carninci, P., Endo, T., Fukuda, S., Fukunishi, Y., Hata, A., Hayatsu, N., Hirozane, T., Horii, F., Ishii, Y., Ishikawa, U., Ishikawa, T., Itoh, M., Izawa, M., Kadota, K., Kagawa, I., Kai, C., Kawai, J., Kikuchi, N., Kiyosawa, H., Kojima, Y., Konno, S., Koya, S., Kurihara, C., Kusake, M., Matsuyama, T., Miki, R., Mizuno, Y., Nakamura, M., Oda, H., Okazaki, Y., Ono, T., Owa, C., Saito, H., Sakai, C., Sato, K., Shibata, K., Shibata, Y., Shigemoto, Y., Shingawa, A., Shiraki, T., Sogabe, Y., Sugahara, Y., Suzuki, H., Suzuki, H., Tagawa, A., Takahashi, F., Tomioka, N., Toyota, T., Tsunoda, Y., Watahiki, A., Watanabe, S., Yamamura, T., Yamanaka, I., Yano, R., Yasunishi, A., Yokota, T., Yoshida, K., Yoshiki, A., Yoshino, M., Muramatsu, M. and Hayashizaki, Y. (2000) The Riken Mouse ESTs (Konno, H., et al.) Unpublished (2000) Contact: Yoshinide Hayashizaki

contact: Yoshihide Hayashizaki  
Laboratory for Genome Exploration Research Group, RIKEN Genomic  
Sciences Center(GSC), Yokohama Institute  
The Institute of Physical and Chemical Research (RIKEN)  
1-7-22 Suenho-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan  
Tel: 81-45-503-9222  
Fax: 81-45-503-9216  
Email: genome-res@gsc.riken.go.jp,  
url:http://genome.gsc.riken.go.jp/  
Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki,  
N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.  
Thermosensitization and thermoactivation of the labile enzymes by  
trehalose and its application for the synthesis of full length  
cDNA. *Proc. Natl. Acad. Sci. U.S.A.* 95 (2), 520-524 (1998)  
Itoh,M., Kitsuai,T., Akiyama,T., Shibata,K., Izawa,M., Kawai,J.,  
Tomaru,Y., Carninci,P., Shibata,Y., Ozawa,Y., Muramatsu,M., Okazaki,  
Y. and Hayashizaki,Y.  
Automated filtration-based high-throughput plasmid preparation  
system. *Genome Res.* 9 (5), 463-470 (1999)  
Carninci,P. and Hayashizaki,Y.  
High-efficiency full-length cDNA cloning. *Methods Enzymol.* 303,  
19-44 (1999)  
Please visit our web site (<http://genome.rtc.riken.go.jp>) for  
further details.

Location/Qualifiers  
1. .224

```

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="4930563D02"
/clone_1fb="RIKEN full-length enriched, adult male testis
(DH10B)"
/sex="male"
/tissue_type="testis"
/dev_stage="adult"
/lab_host="DH10B"
/notes="Site_1: SalI; Site_2: BamHI. cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
plimed with a primer [5',
GAGGAGAGAGAGCATCCACAGAGCTCTTTTTTTTTTTTTTTNN 3'], cDNA was
prepared by using triethanol thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. Second strand cDNA was prepared with the
primer adapter of sequence [5',
GAGGAGAGAGATTCGATTCGATTAAATTAATATCCTCCCCCCCCCCC 3']. cDNA
was cloned into the XhoI and BamHI sites. Vector: a
modified pluescript KS(+) after bulk excision from Lambda
FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI."

```

85.5%; Score 18.8; DB 10; Length 224;

```

Qy 1 AAGAAAAATCTAGACAAGCAA 22
    |||||  |||||
Db 160 AAGAAAAATTCAGACAAGCAA 181

```

LOCUS	424 bp	DNA	linear
BH670801			GSS-19-FEB-2007
DEFINITION	BOMK125TF	BO_2_3_KB	Brassicaceae clemathea genomic clone BOMK125, DNA sequence.
ACCESSION	BH670801		
VERSION	BH670801.1		GI:18734592

**Brassica oleracea**

REFERENCE	1 (bases 1 to 424)
AUTHORS	Tow, C.D., Van Aken, S., Utterback, T., and Fraser, C.M.
TITLE	Whole genome shotgun sequencing of <i>Brassica oleracea</i>
JOURNAL	Unpublished (2001)
COMMENT	Contact: Chris Towm

Location/Qualifiers  
1. .424

```

/organism="Brassicaceae"
/strain="TO1000DH3"
/db_xref="taxon:3712"
/clone="BOMK25"
/clone_1lb="BO_2_3-KB"
/notes="Vector: pHS01, site_1: BstXI, 2-3 kb sheared
genomic DNA inserted into pHS01 using BstXI linkers"
BASE COUNT
165 a      86 c      66 g      107 t
ORIGIN

```

Query Match	85.58;	Score 18.8;	DB 17;	Length 424;
Best Local Similarity	90.9%;	Pred. No. 3.8e+03;		
Matches 20; Conservative	0;	Mismatches 2;	Indels 0;	Gaps 0;

```

QY      1  AAGAAAAAATCTAGACCAAGCAA  22
          |||| ||||| |||||
Db      127 AAGATAAAATCTAGATAAGCAA  148

```

RESULT	6
LOCUS	AZ009207/c
DEFINITION	RPCI-23-365C18.TV RPCI-23 Mus musculus genomic clone RPCI-23-365C18 DNA sequence.
ACCESSION	AZ009207
VERSION	AZ009207.1
KEYWORDS	GSS.
SOURCE	house mouse.

REFERENCE  
1 (bases 1 to 449)  
Zhao, S., Niernan, W., Feldblum, T., Malek, J., Shatsman, S., Aktenret  
AUTHORS

TITLE  
Journal  
Mouse BAC End Sequences from Library RPCI-23  
Unpublished (1999)

## COMMENT

Other GSSs: RPCI-23-365C18.TJ  
 Contact: Shaying Zhao  
 Department of Eukaryotic Genomics  
 The Institute for Genomic Research  
 9712 Medical Center Dr., Rockville, MD 20850, USA  
 Tel: 301 838 0200  
 Fax: 301 838 0200

Email: szhaoc@tigr.org  
 Clones are derived from the mouse BAC library RPCI-23. For BAC library availability, please contact Pieter de Jong (pieter@edjlong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/orderingframe.html) or from Resea ch Genetics (info@resgen.com). BAC end page: http://www.tigr.org/tdb/bac\_ends/mouse/bac\_end\_intro.html  
 Plate: 365 row: C column: 18  
 Seq primer: T7  
 Class: BAC ends.

## FEATURES

## source

## Location/Qualifiers

1..449  
 /organism="Mus musculus"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="RPCI-23-365C18"  
 /clone\_lib="RPCI-23"  
 /sex="Female"  
 /lab\_host="DH10B"  
 /note="Organ: Kidney/Brain; Vector: pBAC3.6; Site: 1; EORI: Site-2; EORI: Female C57BL/6J mouse kidney and/or brain genomic DNA was isolated and partially digested with a combination of EORI and EORI Methylase. Size selected DNA was cloned into the pBAC3.6 vector at the EORI sites. The ligation products were transformed into DH10B electrocompetent cells (BRL Life Technologies)."

## BASE COUNT

97 a 79 c 80 g 193 t

## ORIGIN

Query Match 85.5%; Score 18.8; DB 17; Length 449;  
 Best Local Similarity 90.9%; Pred. No. 3.8e+03;  
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTAGACACGCA 22

Db 423 AATGAAAAATCTAGACACGCA 402

## RESULT 7

## LOCUS

BB768795 476 bp mRNA linear EST 17-OCT-2001  
 BB768795 RIKEN full-length enriched, B16 F10Y cells Mus musculus

## DEFINITION

cDNA clone G370089D24 3', mRNA sequence.

## ACCESSION

BB768795

## VERSION

BB768795.1 GI:16211337

## KEYWORDS

EST.

## SOURCE

house mouse.

## ORGANISM

Mus musculus; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Eukaryota; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

## REFERENCE

1 (bases 1 to 476)

## AUTHORS

Akimura,T., Arakawa,T., Carninci,P., Furuno,M., Hanagaki,T., Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirazawa,T., Imoto,T., Ito,M., Kawai,J., Kojima,Y., Komori,H., Kouda,M., Matsuyama,T., Nakamura,M., Nishi,K., Nomura,K., Numasaki,R., Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sakazume,N., Sasaki,D., Sato,K., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagawa,A., Takahashi,F., Takaku-Akhirita,S., Tanaka,T., Tomaru,A., Toya,T., Watahiki,A., Yasunishi,A., Muramatsu,M. and Hayashizaki,Y.  
 RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura,T., et al. 2001)

## TITLE

Unpublished (2001)

## JOURNAL

Contact: Yoshinobu Hayashizaki  
 Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute  
 The Institute of Physical and Chemical Research (RIKEN)

## FEATURES

## source

## Location/Qualifiers

1..476  
 /organism="Mus musculus"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="G370089D24"  
 /clone\_lib="RIKEN full-length enriched, B16 F10Y cells"  
 /cell\_type="B16 F10Y cells"  
 /note="pooled tissues; (tissue\_type=cerebellum, dev\_stage=16 days neonate, sex=mixed), (tissue\_type=cerebellum, dev\_stage=0 day neonate, sex=mixed), (tissue\_type=hippocampus, dev\_stage=adult, sex=male), (tissue\_type=whole body, dev\_stage=9 days embryo, sex=mixed), (tissue\_type=lung, dev\_stage=13 days embryo, sex=mixed)"  
 e mouse tissues.

## BASE COUNT

128 a 105 c 111 g 132 t

## ORIGIN

Query Match 85.5%; Score 18.8; DB 10; Length 476;  
 Best Local Similarity 90.9%; Pred. No. 3.8e+03;  
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTAGACACGCA 22

Db 303 AAAAAAAACCTAGACACGCA 282

## RESULT 8

## LOCUS

BF004802 505 bp mRNA linear EST 06-OCT-2000  
 BF004802 KVL Medicago truncatula cDNA clone pKV1-18B14, mRNA

## DEFINITION

sequence.

## ACCESSION

BF004802

## VERSION

BF004802.1 GI:10705077

## KEYWORDS

EST.

## SOURCE

barrel medic.

## ORGANISM

Medicago truncatula

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliopsida; eudicotyledons; core eudicots; Rosidae; eustosids I; Fabales; Fabaceae; Papilionoideae; Trifoliaceae; Medicago.

## REFERENCE

1 (bases 1 to 505)

## AUTHORS

Vandenbosch,K., Endre,G., Hur,J., Moore,J., Beremand,P., Ellis,L., Town,C.D., Bowman,C.L., Craven,M.B., Hansen,T.S., Holt,L.E. and Fraser,C.M.  
 ESTs from roots of Medicago truncatula 24 hours after inoculation with Sinorhizobium meliloti  
 Unpublished (1999)

## TITLE

Unpublished (1999)

## JOURNAL

## COMMENT

Contact: VandenBosch K  
Department of Plant Biology  
University of Minnesota  
220 Biosci Center, 1445 Gortner Ave, St. Paul, MN 55108, USA  
Tel: 612 624 2755  
Fax: 612 625 1738  
Email: kvandenbosch.umn.edu  
Texas A&M University name: T268612e TIGR sequence name: MTIBJ077K  
More information is available at: <http://chrysie.tamu.edu/medicago>  
Seq primer: SKmod (CTA GAA CTA gty gat CC).  
Location/Qualifiers

## FEATURES

## source

1. 505  
/organism="Medicago truncatula"  
/cultivar="genotype A17"  
/db\_xref="taxon:3880"  
/clone="PKV1-18B14"  
/clone\_1lb="KV1"  
/issue\_type="Seedling roots"  
/dev\_stage="24 hours post-inoculation with Sinorhizobium  
meliloti"  
/lab\_host="E. coli strain XLOLR"  
/note="Vector: pBluescript SK-; Site\_1: EcoRI; Site\_2:  
XhoI; cDNA was prepared from polyA+ enriched RNA. The cDNA  
was directionally ligated into the unzip XR vector from  
stratagene and packaged using gigapack iii gold packaging  
extracts. Plasmids containing cDNA inserts were excised  
from the recombinant lambda-Zap phage using Ex-assist  
helper phage and propagated in XLOLR cells."  
BASE COUNT 144 a 93 c 103 g 164 t 1 others  
ORIGIN

## Query Match

Best Local Similarity 90.9%; Score 18.8; DB 12; Length 505;  
Pred. No. 3.7e+03;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTGACACAGCAA 22  
||||| ||||| ||||| |||||

Db 109 AAGAAAAATCTGACACAGCAA 88

RESULT 9 AA447445 528 bp mRNA linear EST 04-JUN-1997  
LOCUS zw93g12.r1 Soares\_total.fetus\_Nb2HF8\_9w Homo sapiens cDNA clone

DEFINITION IMAGE:784582 5', mRNA sequence.  
AA447445  
VERSION AA447445.1 GI:2161115

KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
1 (bases 1 to 528)

AUTHORS Hillier, L., Allen, M., Bowles, L., Dubouche, T., Giesel, G., Jost, S.,  
Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B.,  
Schellenberg, K., Stepien, M., Tan, F., Theising, B., White, Y., Wyllie,  
T., Waterston, R. and Wilson, R.  
WashU-Merck EST Project 1997

TITLE Unpublished (1997)  
JOURNAL Contact: Wilson RK  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810

FEATURES  
source  
1. 528  
/organism="Homo sapiens"  
/db\_xref="GDB:5982200"  
Location/Qualifiers

/db\_xref="taxon:9606"  
/clone="IMAGE:784582"  
/clone\_1lb="Soares\_total.fetus\_Nb2HF8\_9w"  
/dev\_stage="8-9 weeks"  
/lab\_host="DH10B"  
/note="Vector: pT73D-Pac (Pharmacia) with a modified  
polylinker; Site\_1: Not I; Site\_2: Eco RI; 1st strand cDNA  
was prepared from mRNA obtained from pooled 8-9 week  
(total) fetus material with a Not I - oligo(dT) primer [5'  
GTGACCAATCTGAAGTGGAGCGCCCTTAATTTTCTTTTCTTTT 3'].  
Double-stranded cDNA was ligated to Eco RI adaptors  
(Pharmacia), digested with Not I and cloned into the Not I  
and Eco RI sites of the modified pT73 vector. Library  
went through one round of normalization, and was  
constructed by Benito Soares and M. Fatima Bonaldo."

BASE COUNT 155 a 113 c 91 g 169 t  
ORIGIN

Query Match 85.5%; Score 18.8; DB 9; Length 528;  
Best Local Similarity 90.9%; Pred. No. 3.7e+03;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGAAAAATCTGACACAGCAA 22  
||||| ||||| ||||| |||||

Db 24 AAGAAAAATCTGACACAGCAA 45

RESULT 10 A0414714 544 bp DNA linear GSS 23-MAR-1999  
LOCUS RPCI-11-171019.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-171019

DEFINITION RPCI-11-171019.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-171019  
DNA sequence.  
A0414714  
VERSION A0414714.1 GI:4473683

KEYWORDS GSS.  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
1 (bases 1 to 544)

AUTHORS Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter,  
J.C.  
Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready  
Map Building  
Unpublished (1997)

JOURNAL Contact: Shaying Zhao, William Nierman, Mark Adams  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850  
Tel: 301 838 0200  
Fax: 301 838 0208

TITLE Email: hbeetlgr.org  
Clones are derived from the human BAC library RPCI-11. For BAC  
library availability, please contact Pieter de Jong  
(pieter@dejong.med.buffalo.edu). Clones may be purchased from  
BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from  
Research Genet cs ([inforesgen.com](http://inforesgen.com)). BAC end search page:  
[http://www.tigr.org/tdb/humgen/bac\\_end\\_search/bac\\_end\\_search.html](http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html).  
Seq primer: SP6  
Class: BAC ends.

FEATURES  
source  
1. 544  
Location/Qualifiers

/organism="Homo sapiens"  
/db\_xref="GDB:7565634"  
/db\_xref="taxon:9606"  
/clone="RPCI-11-171019"  
/clone\_1lb="RPCI-11"  
/sex="Male"  
/cell\_type="Lymphocytes"  
/note="Vector: pBAC6.6; Site\_1: EcoRI; Site\_2: EcoRI;  
RPCI11 Human Male BAC Library"  
BASE COUNT 189 a 97 c 100 g 158 t  
ORIGIN

[illegible]

DEFINITION	RPCT-11-353p23.tv RPCT-11 Homo sapiens genomic clone RPCT-11-353p23				
ACCESSION	RPCT-11-353p23.DNA sequence.				
VERSION	A0534621				
KEYWORDS	A0534621.1 GI:4846311				
SOURCE	GSS.				
ORGANISM	human.				
REFERENCE	Homo sapiens				
AUTHORS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
TITLE	1 (bases 1 to 567)				
JOURNAL	Zhao,S., Adams,M.D., Nierman,W., Malek,J., de Jong,P. and Venter,J.C.				
COMMENT	Use of BAC End Sequences from Library RPCT-11 for Sequence-Ready Map Building				
	Unpublished (1997)				
	Other-GSSs: RPCT-11-353p23.TV				
	Contact: Shaying Zhao, William Nierman, Mark Adams				
	Department of Eukaryotic Genomics				
	The Institute for Genomic Research				
	9712 Medical Center Dr., Rockville, MD 20850				
	Tel: 301 838 0200				
	Fax: 301 838 0208				
	Email: hbe@tigr.org				
	Clones are derived from the human BAC library RPCT-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources ( <a href="http://bacpac.med.buffalo.edu/ordering">http://bacpac.med.buffalo.edu/ordering</a> ) or from Research Genet cs ( <a href="http://inforesgen.com">inforesgen.com</a> ). BAC end search page: <a href="http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html">http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html</a> .				
	Seq primer: T7				
	Class: BAC ends.				
FEATURES	location/Qualifiers				
source	1..567				
	/organism="Homo sapiens"				
	/db_xref="GDB:7635550"				
	/db_xref="taxon:9606"				
	/clone="RPCT-11-353p23"				
	/clone_1db="RPCT-11"				
	/sex="Male"				
	/cell_type="Lymphocytes"				
	/note=Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI; RPCT11 Human Male BAC Library"				
BASE COUNT	191 a 99 c 111 g 165 t 1 others				
ORIGIN					
Query Match	85.5%; Score 18.8; DB 17; Length 567;				
Best Local Similarity	90.9%; Pred. No. 3.7e+03;				
Matches	20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;				
OY	1 AAGAAAAATCTAGACAAAGCAA 22				
Db	476 AAGAAAAAGCTAGACAAAGAA 497				
RESULT 13					
LOCUS	A2506877 568 bp DNA linear GSS 05-OCT-2000				
DEFINITION	IM0348C14F Mouse 10kb plasmid UUGC1M library Mus musculus genomic				
ACCESSION	clone UUGC1M0348C14 F, DNA sequence.				
VERSION	A2506877				
KEYWORDS	A2506877.1 GI:10668193				
SOURCE	GSS.				
ORGANISM	house mouse.				
	Mus musculus				
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
REFERENCE	1 (bases 1 to 568)				
AUTHORS	Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meeren,E., Pedersen,T., Rellly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D.,Weiss,R.				
TITLE	Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts				

JOURNAL  
COMMENT

Unpublished (2000)  
Contact: Robert B. Weiss  
University of Utah Genome Center  
University of Utah  
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT  
84112, USA  
Tel: 801 585 5606  
Fax: 801 585 7177  
Email: ddunne@genetics.utah.edu  
Insert length: 10000 Std Error: 0.00  
Plate: 0348 Row: C Column: 14  
Seq primer: CTTGTAAACGACGCCACAGT  
Class: plasmid ends  
High quality sequence stop: 568.

## FEATURES

## source

1. 568  
Location/Qualifiers  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UUC1M0348C14"  
/clone\_lib="Mouse 10kb plasmid UUC1M library"  
/sex="Male"  
/lab\_host="E. coli strain XL10-Gold, T1-resistant, F-"  
/note="Vector: FMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource  
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g1473211419b1AF129072.1), a copy number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT  
ORIGIN

192 a 101 c 123 g 150 t 2 others  
Query Match 85.5%; Score 18.8; DB 17; Length 568;  
Best Local Similarity 90.9%; Pred. No. 3.7e+03;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

1 AAGAAAAATCTAGACACGAA 22  
|||||  
Db 191 AAGAAAAATCTAGACACGAA 212

RESULT 14  
LOCUS

DEFINITION A0534617 597 bp DNA linear GSS 18-MAY-1999  
R00111-353P21.TV R00111 Homo sapiens genomic clone R00111-353P21  
DNA sequence.

ACCESSION A0534617  
VERSION A0534617.1 GI:4846307  
KEYWORDS GSS.  
SOURCE human.  
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 597)  
Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter, J.C.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

TITLE Use of BAC End Sequences from Library R00111 for Sequence-Ready Map Building  
JOURNAL Unpublished (1997)  
COMMENT Other\_GSSs: R00111-353P21.TV

Contact: Shaying Zhao, William Nierman, Mark Adams  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850  
Tel: 301 838 0200  
Fax: 301 838 0208  
Email: hbe@tigr.org

Clones are derived from the human BAC library R00111. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/tldb/humgen/bac\_end\_search/bac\_end\_search.html.  
Seq primer: T7  
Class: BAC ends.

## FEATURES

## source

1. 597  
Location/Qualifiers  
/organism="Homo sapiens"  
/db\_xref="GDB:7635548"  
/db\_xref="taxon:9606"  
/clone="R00111-353P21"  
/clone\_lib="R00111"  
/sex="Male"  
/cell\_type="Lymphocytes"  
/note="Vector: pBAC3.6; Site\_1: EcoRI; Site\_2: EcoRI; R00111 Human Male Library"

BASE COUNT  
ORIGIN

201 a 107 c 115 g 174 t  
Query Match 85.5%; Score 18.8; DB 17; Length 597;  
Best Local Similarity 90.9%; Pred. No. 3.6e+03;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

1 AAGAAAAATCTAGACACGAA 22  
|||||  
Db 496 AAGAAAAATCTAGACACGAA 517

RESULT 15  
LOCUS

DEFINITION BH552202 710 bp DNA linear GSS 14-DEC-2001  
BOH0N17TF BOH0 Brassica oleracea genomic clone BOH0N17, DNA sequence.

ACCESSION BH552202  
VERSION BH552202.1 GI:17803982  
KEYWORDS GSS.  
SOURCE Brassica oleracea.  
ORGANISM Brassica oleracea

REFERENCE 1 (bases 1 to 710)  
Town, C.D., Van Aken, S., Utterback, T. and Fraser, C.M.  
Whole genome shotgun sequencing of Brassica oleracea  
Unpublished (2001)  
Other\_GSSs: BOH0N17TF  
Contact: Chris Town  
TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA.  
Tel: 301-838-9523  
Fax: 301-838-0208  
Email: cdtown@tigr.org  
DNA is from a doubled haploid provided by Tom Osborn.  
Seq primer: T7  
Class: sheared ends.

## FEATURES

## source

1. 710  
Location/Qualifiers  
/organism="Brassica oleracea"  
/strain="TO1000DH3"  
/db\_xref="taxon:3712"  
/clone="BOH0N17"  
/clone\_lib="BOH0"  
/note="Vector: pBAC3.6; Site\_1: BstXI; 2-3 kb sheared

BASE COUNT            genomic DNA inserted into PHOS1 using BstXI linkers"  
ORIGIN            209 a    146 c    149 g    206 t

Query Match            85.5%; Score 18.8; DB 17; Length 710;  
Best Local Similarity 90.9%; Pred. No. 3.5e+03;  
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY    1 AAGAAAAATCTAGACAGCAA 22  
      |||| |||||||| |||||  
Db    154 AAGATTAATCTAGATAAGCAA 133

Search completed: March 17, 2003, 13:09:20  
Job time : 782.688 secs



GenCore version 5.1.4-p5\_4578  
Copyright (c) 1993 - 2003 Compugen Ltd.

## OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 09:43:42 ; Search time 230.108 Seconds  
(without alignments)

3161.870 Million cell updates/sec

Title: US-09-836-439-5

Perfect score: 25  
Sequence: 1 gcttcttgcgtcagagctctcca 25

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

## Database :

GenEmbl:\*

- 1: gb\_da:\*
- 2: gb\_hlg:\*
- 3: gb\_in:\*
- 4: gb\_cm:\*
- 5: gb\_ov:\*
- 6: gb\_pat:\*
- 7: gb\_ph:\*
- 8: gb\_pl:\*
- 9: gb\_pr:\*
- 10: gb\_pro:\*
- 11: gb\_scs:\*
- 12: gb\_sy:\*
- 13: gb\_un:\*
- 14: gb\_vl:\*
- 15: em\_ba:\*
- 16: em\_fun:\*
- 17: em\_hum:\*
- 18: em\_in:\*
- 19: em\_mu:\*
- 20: em\_cm:\*
- 21: em\_or:\*
- 22: em\_ov:\*
- 23: em\_pat:\*
- 24: em\_ph:\*
- 25: em\_pl:\*
- 26: em\_ro:\*
- 27: em\_scs:\*
- 28: em\_un:\*
- 29: em\_vl:\*
- 30: em\_hlg\_hum:\*
- 31: em\_hlg\_inv:\*
- 32: em\_hlg\_other:\*
- 33: em\_hlg\_mus:\*
- 34: em\_hlg\_pln:\*
- 35: em\_hlg\_rtd:\*
- 36: em\_hlg\_mam:\*
- 37: em\_hlg\_vrt:\*
- 38: em\_sy:\*
- 39: em\_hlg\_hum:\*
- 40: em\_hlg\_mus:\*
- 41: em\_hlg\_other:\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	23.4	93.6	269	10 MUSOPSA4	M36698 Mouse opsin
2	23.4	93.6	2610	10 BC031766	BC031766 Mus muscu
3	23.4	93.6	3249	10 BC013125	BC013125 Mus muscu
4	23.4	93.6	9483	10 MUSOPS	M55171 Mouse opsin
5	21.8	87.2	175	5 AF249155	AF249155 Indirana
6	21.8	87.2	1062	5 TN1293018	AJ295018 Tetradon
7	20.4	81.6	178639	2 AC095437	AC095437 Rattus no
8	20.2	80.8	169	5 AF249131	AF249131 Microhya
9	20.2	80.8	175	5 AF249129	AF249129 Bufo mela
10	20.2	80.8	175	5 AF249133	AF249133 Mantella
11	20.2	80.8	175	5 AF249134	AF249134 Mantidact
12	20.2	80.8	175	5 AF249136	AF249136 Boophis x
13	20.2	80.8	175	5 AF249137	AF249137 Boophis t
14	20.2	80.8	175	5 AF249138	AF249138 Laliostom
15	20.2	80.8	175	5 AF249139	AF249139 Referyary
16	20.2	80.8	175	5 AF249140	AF249140 Referyary
17	20.2	80.8	175	5 AF249141	AF249141 Hoplobatr
18	20.2	80.8	175	5 AF249142	AF249142 Sphaerotr
19	20.2	80.8	175	5 AF249143	AF249143 Euphycti
20	20.2	80.8	175	5 AF249144	AF249144 Nannophry
21	20.2	80.8	175	5 AF249146	AF249146 Nyctibatr
22	20.2	80.8	175	5 AF249147	AF249147 Limnodyn
23	20.2	80.8	175	5 AF249148	AF249148 Limnodyn
24	20.2	80.8	175	5 AF249149	AF249149 Rana curt
25	20.2	80.8	175	5 AF249150	AF249150 Rana temp
26	20.2	80.8	175	5 AF249151	AF249151 Rana temp
27	20.2	80.8	175	5 AF249152	AF249152 Microxalu
28	20.2	80.8	175	5 AF249154	AF249154 Indirana
29	20.2	80.8	175	5 AF249156	AF249156 Polypedat
30	20.2	80.8	175	5 AF249157	AF249157 Rhacophor
31	20.2	80.8	175	5 AF249158	AF249158 Philautus
32	20.2	80.8	405	5 AF221974	AF221974 Bufo pusi
33	20.2	80.8	416	5 AF221979	AF221979 Bufo pant
34	20.2	80.8	825	5 AF137213	AF137213 Ostracion
35	20.2	80.8	924	5 AB084931	AB084931 Dimidioc
36	20.2	80.8	924	5 AB084938	AB084938 Oreochrom
37	20.2	80.8	924	5 AB084940	AB084940 Sarothero
38	20.2	80.8	924	5 AB084941	AB084941 Spatiodus
39	20.2	80.8	924	5 AB084944	AB084944 Tilapia r
40	20.2	80.8	924	5 AB084947	AB084947 Xenotilap
41	20.2	80.8	1053	5 AF021242	AF021242 Melopsitt
42	20.2	80.8	1059	5 AMU12328	U12328 Asitynax me
43	20.2	80.8	1399	5 RTU59920	RTU59920 Rana temp
44	20.2	80.8	1423	5 MSU118666	MSU118666 Mullus surm
45	20.2	80.8	1469	5 MVU57539	MVU57539 Myrtilistis

## ALIGNMENTS

RESULT 1	MUSOPSA4	269 bp	DNA	Linear	ROD 08-MAY-1993
LOCUS	MUSOPSA4				
DEFINITION	Mouse opsin gene, exon 4.				
ACCESSION	M36698 X69175				
VERSION	M36698.1 GI:200149				
KEYWORDS	opsin.				
SEGMENT	4 of 5				
SOURCE	Mus musculus (strain C56BL/6J) eye DNA.				
ORGANISM	Mus musculus				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
AUTHORS	Baehr, W., Falk, J.D., Bugra, K., Triantafyllou, J.T. and McGinnis, J.F.				
TITLE	Isolation and analysis of the mouse opsin gene				

JOURNAL FEB8 lett. 238 (2), 253-256 (1988)  
MEDLINE 89005694  
PUBMED 2844600

FEATURES  
source Location/Qualifiers

1..269  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/tissue\_type="eye"  
order(M36897.1:170, >189,<1, .14)  
/gene="Opsin"  
/note="117 bp gap"  
/number=3  
15..254  
/gene="Opsin"  
/number=4  
exon

BASE COUNT 57 a 87 c 58 g 67 t  
ORIGIN About 117 bp after segment 3.

Query Match 93.6%; Score 23.4; DB 10; Length 269;  
Best Local Similarity 96.0%; Pred. No. 1.4;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTTGCTGAGAGCTCTTCCA 25  
|||||  
Db 192 GCTTCTTTGCTGAGAGCTCTTCCA 216

RESULT 2  
BC031766 2610 bp mRNA linear ROD 07-AUG-2002  
LOCUS Mus musculus, clone MGC:25387 IMAGE:4527040, mRNA, complete cds.  
DEFINITION BC031766  
ACCESSION BC031766.1 GI:21594394  
KEYWORDS MGC.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE  
AUTHORS Strausberg, R.  
TITLE Direct Submission  
JOURNAL Submitted (06-JUN-2002) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA

REMARK  
COMMENT NIH-MGC Project URL: <http://mgc.nci.nih.gov>  
Contact: MGC help desk  
Email: [cgabs-remail.nih.gov](mailto:cgabs-remail.nih.gov)  
Tissue Procurement: The Cepko Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Baylor College of Medicine Human Genome Sequencing Center  
Center code: BCM-HGSC  
Web site: <http://www.hgsc.bcm.tmc.edu/cdna/>  
Contact: [amg@bcm.tmc.edu](mailto:amg@bcm.tmc.edu)  
Gunnarntne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Hale, S.M., Yoon, V.S., Kowis, C.R., Lawrence, S., Martin, R.G., Muzny, D.M., Richards, S., Gibbs, R.A.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
Series: IRAC Plate: 31 Row: n Column: 2  
This clone was selected for full length sequencing because it passed the following selection criteria: Hexamer frequency ORF analysis, GenomScan gene prediction.  
Location/Qualifiers

FEATURES  
source 1..2610  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="MGC:25387 IMAGE:4527040"  
/tissue\_type="Eye, retina, mouse strain C57Bl\6"

/clone\_11b="NIH\_MGC\_94"  
/lab\_host="DH10B"  
/note="Vector: PCMV-SPORT6"  
80..1126  
/product="Unknown (protein for MGC:25387)"  
/protein\_id="AAH31766.1"  
/db\_xref="GI:21594395"  
/db\_xref="locustid:212541"  
/translation="MNGTEGPNEVYFESNVTVGVSPEEPQYLLAEPMQSMIAAYM  
FLIIVGLFPIINFLIVYVQHKRLPLNYILNLAVDLFWFGFTTITSLHG  
FVGEPTCNLEGFATLGEELIALSVLIVYVAVVCKPNSFEGNHAIWVET  
WMLALCAAPPLVGMRSRYIPBMGSCGIDYVTLKPEVNSEFVLYMVFHTTIPMIV  
IPFCYGLVFTYKRAAQQQESATTDKAKETRYIIVIVIEFLICMIPYASVARYIF  
THGGSNFGPIFMTLPAPFAKSSSIINPVIYIMLNQFRMCMITLCCGKNPLGDDAS  
ATASKTETSQVAPA"

BASE COUNT 604 a 727 c 656 g 623 t  
ORIGIN

Query Match 93.6%; Score 23.4; DB 10; Length 2610;  
Best Local Similarity 96.0%; Pred. No. 1.5;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTTGCTGAGAGCTCTTCCA 25  
|||||  
Db 953 GCTTCTTTGCTGAGAGCTCTTCCA 977

RESULT 3  
BC013125 3249 bp mRNA linear ROD 07-AUG-2002  
LOCUS Mus musculus, similar to rhodopsin (opsin 2, rod pigment) (retinitis pigmentosa 4, autosomal dominant), clone MGC:21565  
DEFINITION BC013125  
ACCESSION BC013125.1 GI:15341884  
KEYWORDS MGC.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE  
AUTHORS Strausberg, R.  
TITLE Direct Submission  
JOURNAL Submitted (27-AUG-2001) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA

REMARK  
COMMENT NIH-MGC Project URL: <http://mgc.nci.nih.gov>  
Contact: MGC help desk  
Email: [cgabs-remail.nih.gov](mailto:cgabs-remail.nih.gov)  
Tissue Procurement: The Cepko Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Baylor College of Medicine Human Genome Sequencing Center  
Center code: BCM-HGSC  
Web site: <http://www.hgsc.bcm.tmc.edu/cdna/>  
Contact: [amg@bcm.tmc.edu](mailto:amg@bcm.tmc.edu)  
Gunnarntne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Hale, S.M., Yoon, V.S., Kowis, C.R., Lawrence, S., Martin, R.G., Muzny, D.M., Richards, S., Gibbs, R.A.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
Series: IRAC Plate: 28 Row: e Column: 21  
This clone was selected for full length sequencing because it passed the following selection criteria: Similarity but not identity to protein.  
Location/Qualifiers

FEATURES  
source 1..3249  
/organism="Mus musculus"  
/db\_xref="taxon:10090"





TITLE  
JOURNAL  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

Direct Submission  
Unpublished  
2 (bases 1 to 178639)  
Worley, K.C.  
Direct Submission  
Submitted (17-SEP-2001) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA  
3 (bases 1 to 178639)  
Worley, K.C.  
Direct Submission  
Submitted (10-JUL-2002) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA  
On Jul 9, 2002 this sequence version replaced gi:17941914.

----- Genome Center  
Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: hgsc-help@bcm.tmc.edu

----- Project Information  
Center project name: CGHA  
Center clone name: CH230-4H9

----- Summary Statistics  
Sequencing vector: Plasmid  
Chemistry: Dye-terminator Big Dye: 100% of reads  
Assembly program: Phrap: version 0.990329  
Consensus quality: 128571 bases at least Q40  
Consensus quality: 133786 bases at least Q30  
Consensus quality: 137554 bases at least Q20

----- NOTE: Estimated insert size may differ from sequence length  
(see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_drift\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_drift_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 67 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1  
1037 1036: contig of 1036 bp in length  
1137 1136: gap of unknown length  
2378 2377: contig of 1241 bp in length  
2478 2477: gap of unknown length  
3635 3634: contig of 1157 bp in length  
3735 3734: gap of unknown length  
5028 5027: contig of 1293 bp in length  
5128 5127: gap of unknown length  
6328 6327: contig of 1200 bp in length  
6428 6427: gap of unknown length  
8035 8034: contig of 1607 bp in length  
8135 8134: gap of unknown length  
9425 9424: contig of 1290 bp in length  
9525 9524: gap of unknown length  
10939 10938: contig of 1414 bp in length  
11039 11038: gap of unknown length  
12078 12077: contig of 1039 bp in length  
12178 12177: gap of unknown length  
13191 13190: contig of 1013 bp in length  
13291 13290: gap of unknown length  
14395 14394: contig of 1104 bp in length  
14495 14494: gap of unknown length  
15667 15666: contig of 1172 bp in length  
15767 15766: gap of unknown length  
17029 17028: contig of 1263 bp in length  
17130 17129: gap of unknown length  
18245 18244: contig of 1115 bp in length  
18345 18344: gap of unknown length  
19988 19987: contig of 1643 bp in length  
20088 20087: gap of unknown length  
21162 21161: contig of 1074 bp in length  
21261 21261: gap of unknown length

21262 22935: contig of 1674 bp in length  
22936 23035: gap of unknown length  
23036 24726: contig of 1691 bp in length  
24727 24826: gap of unknown length  
24827 25984: contig of 1158 bp in length  
25985 26084: gap of unknown length  
26085 28117: contig of 2033 bp in length  
28118 28217: gap of unknown length  
28218 29226: contig of 1009 bp in length  
29227 29326: gap of unknown length  
29327 31330: contig of 2004 bp in length  
31331 31430: gap of unknown length  
31431 33008: contig of 1578 bp in length  
33009 33108: gap of unknown length  
33109 34893: contig of 1785 bp in length  
34894 34993: gap of unknown length  
34994 36418: contig of 1425 bp in length  
36419 38755: gap of unknown length  
38756 38855: contig of 2237 bp in length  
38856 40715: gap of unknown length  
40716 40815: gap of unknown length  
40816 42591: contig of 1776 bp in length  
42592 42691: gap of unknown length  
42692 44432: contig of 1741 bp in length  
44433 44532: gap of unknown length  
44534 46670: contig of 2138 bp in length  
46671 46770: gap of unknown length  
46771 48171: contig of 1401 bp in length  
48172 48271: gap of unknown length  
48272 50293: contig of 2022 bp in length  
50294 50393: gap of unknown length  
50394 52099: contig of 1706 bp in length  
52100 52199: gap of unknown length  
52200 54742: contig of 2543 bp in length  
54743 54842: gap of unknown length  
54843 57839: contig of 2997 bp in length  
57840 57939: gap of unknown length  
57940 59988: contig of 2050 bp in length  
59989 60089: gap of unknown length  
60090 61699: contig of 1510 bp in length  
61699 61600: gap of unknown length  
61700 63235: contig of 1536 bp in length  
63236 63335: gap of unknown length  
63336 65866: contig of 2361 bp in length  
65867 65796: gap of unknown length  
65797 68230: contig of 2434 bp in length  
68231 68330: gap of unknown length  
68331 70088: contig of 1758 bp in length  
70089 70188: gap of unknown length  
70189 72866: contig of 2678 bp in length  
72867 72966: gap of unknown length  
72967 76485: contig of 3519 bp in length  
76486 76585: gap of unknown length  
76586 78904: contig of 2319 bp in length  
78905 79004: gap of unknown length  
79006 81763: contig of 2755 bp in length  
81764 81863: gap of unknown length  
81864 85381: contig of 3518 bp in length  
85382 85481: gap of unknown length  
85482 88489: contig of 3008 bp in length  
88490 88589: gap of unknown length  
88590 91317: contig of 2728 bp in length  
91318 91417: gap of unknown length  
91418 93443: contig of 2026 bp in length  
93444 95582: contig of 2039 bp in length  
95583 95682: gap of unknown length  
95683 99140: contig of 3458 bp in length  
99141 99240: gap of unknown length  
99241 101817: contig of 2577 bp in length  
101818 101917: gap of unknown length  
101918 105395: contig of 3478 bp in length



Query Match 80.8%; Score 20.2; DB 5; Length 175;  
 Best Local Similarity 88.0%; Pred. No. 52;  
 Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GCTTCTTCTGCTGAGAGCTCTTCCA 25  
 Db 140 GCTTCTTCTGCTGAGAGCTCTGCCA 164

RESULT 11  
 AF249134 175 bp DNA linear VRT 17-JAN-2001  
 LOCUS Mantiactylus cf. ulcersus rhodopsin gene, exon 4 and partial cds.  
 DEFINITION  
 ACCESSION AF249134  
 VERSION AF249134.1 GI:12247197  
 KEYWORDS  
 SOURCE Mantiactylus cf. ulcersus.  
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Rhacophoridae;

REFERENCE 1 (bases 1 to 175)  
 Bossuyt, F. and Miliukovitch, M.C.  
 TITLE Convergent Adaptive Radiations in Madagascan and Asian Ranid Frogs  
 AUTHOR

JOURNAL Unpublished  
 REFERENCE 2 (bases 1 to 175)  
 Bossuyt, F. and Miliukovitch, M.C.  
 TITLE Direct Submission  
 AUTHOR  
 JOURNAL Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of Molecular Biology and Medicine, rue Jeener and Brachet 12,  
 Gosselies B-6041, Belgium

FEATURES  
 Source Location/Qualifiers  
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ORIGIN  
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 /db\_xref="taxon:129014"  
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 /product="rhodopsin"

CDS  
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 VPAPFAKSSAIYNP"  
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BASE COUNT 36 a 50 c 36 g 53 t  
 ORIGIN

Query Match 80.8%; Score 20.2; DB 5; Length 175;  
 Best Local Similarity 88.0%; Pred. No. 52;  
 Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GCTTCTTCTGCTGAGAGCTCTTCCA 25  
 Db 140 GCTTCTTCTGCTGAGAGCTCTGCCA 164

RESULT 12  
 AF249136 175 bp DNA linear VRT 17-JAN-2001  
 LOCUS Boophis xerophilus rhodopsin gene, exon 4 and partial cds.  
 DEFINITION  
 ACCESSION AF249136  
 VERSION AF249136.1 GI:12247201  
 KEYWORDS

SOURCE  
 ORGANISM Boophis xerophilus.  
 Boophis xerophilus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Rhacophoridae;  
 Boophis.

REFERENCE 1 (bases 1 to 175)  
 Bossuyt, F. and Miliukovitch, M.C.  
 TITLE Convergent Adaptive Radiations in Madagascan and Asian Ranid Frogs  
 AUTHOR  
 JOURNAL Unpublished  
 REFERENCE 2 (bases 1 to 175)  
 Bossuyt, F. and Miliukovitch, M.C.  
 TITLE Direct Submission  
 AUTHOR  
 JOURNAL Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of Molecular Biology and Medicine, rue Jeener and Brachet 12,  
 Gosselies B-6041, Belgium

FEATURES  
 source Location/Qualifiers  
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BASE COUNT 34 a 55 c 34 g 52 t  
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Query Match 80.8%; Score 20.2; DB 5; Length 175;  
 Best Local Similarity 88.0%; Pred. No. 52;  
 Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GCTTCTTCTGCTGAGAGCTCTTCCA 25  
 Db 140 GCTTCTTCTGCTGAGAGCTCTGCCA 164

RESULT 13  
 AF249137 175 bp DNA linear VRT 17-JAN-2001  
 LOCUS Boophis tephraeomystax rhodopsin gene, exon 4 and partial cds.  
 DEFINITION  
 ACCESSION AF249137  
 VERSION AF249137.1 GI:12247203  
 KEYWORDS  
 SOURCE Boophis tephraeomystax.  
 ORGANISM Boophis tephraeomystax.  
 Boophis tephraeomystax  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Rhacophoridae;  
 Boophis.

REFERENCE 1 (bases 1 to 175)  
 Bossuyt, F. and Miliukovitch, M.C.  
 TITLE Convergent Adaptive Radiations in Madagascan and Asian Ranid Frogs  
 AUTHOR  
 JOURNAL Unpublished  
 REFERENCE 2 (bases 1 to 175)  
 Bossuyt, F. and Miliukovitch, M.C.  
 TITLE Direct Submission  
 AUTHOR  
 JOURNAL Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of Molecular Biology and Medicine, rue Jeener and Brachet 12,  
 Gosselies B-6041, Belgium

FEATURES  
 source Location/Qualifiers  
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ORIGIN  
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 /db\_xref="taxon:68440"  
 <1..>175  
 /product="rhodopsin"

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VPAFFAKSSAIYNP"
<1..>175
exon
/number=4
BASE COUNT      35 a      56 c      33 g      51 t
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Query Match      80.8%; Score 20.2; DB 5; Length 175;
Best Local Similarity 88.0%; Pred. No. 52;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 GCTTCTTGTGAGAGCTCTGCCA 25
|||||
Db 140 GCTTCTTGTGCCAAGAGCTCTGCCA 164

RESULT 14
AF249138      175 bp      DNA      linear      VRT 17-JAN-2001
LOCUS      Laliostoma labrosum rhodopsin gene, exon 4 and partial cds.
DEFINITION
AF249138
ACCESSION
VERSION
AF249138.1 GI:12247205
KEYWORDS
SOURCE      Laliostoma labrosum.
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
      Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Ranidae;
      Laliostoma.
REFERENCE
AUTHORS      1 (bases 1 to 175)
      Bossuyt, F. and Milinkovitch, M.C.
TITLE      Convergent Adaptive Radiations in Madagascar and Asian Ranid Frogs
      Reveal Co-variation between Larval and Adult Traits
JOURNAL
REFERENCE
AUTHORS      2 (bases 1 to 175)
      Bossuyt, F. and Milinkovitch, M.C.
TITLE      Direct Submission
      Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of
      Molecular Biology and Medicine, rue Jeener and Brachet 12,
      Gosselies B-6041, Belgium
FEATURES
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BASE COUNT      35 a      55 c      36 g      49 t
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Best Local Similarity 88.0%; Pred. No. 52;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 GCTTCTTGTGAGAGCTCTGCCA 25
|||||
Db 140 GCTTCTTGTGCCAAGAGCTCTGCCA 164

RESULT 15
AF249139      175 bp      DNA      linear      VRT 17-JAN-2001
LOCUS

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DEFINITION      Fejervarya syhadrensis rhodopsin gene, exon 4 and partial cds.
ACCESSION      AF249139
VERSION      AF249139.1 GI:12247207
KEYWORDS
SOURCE
ORGANISM      Fejervarya syhadrensis.
      Fejervarya syhadrensis
      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
      Amphibia; Batrachia; Anura; Neobatrachia; Ranoidae; Ranidae;
      Fejervarya.
REFERENCE
AUTHORS      1 (bases 1 to 175)
      Bossuyt, F. and Milinkovitch, M.C.
TITLE      Convergent Adaptive Radiations in Madagascar and Asian Ranid Frogs
      Reveal Co-variation between Larval and Adult Traits
JOURNAL
REFERENCE
AUTHORS      2 (bases 1 to 175)
      Bossuyt, F. and Milinkovitch, M.C.
TITLE      Direct Submission
      Submitted (27-MAR-2000) Unit of Evolutionary Genetics, Institute of
      Molecular Biology and Medicine, rue Jeener and Brachet 12,
      Gosselies B-6041, Belgium
FEATURES
source
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      <1..>175
      /product="rhodopsin"
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BASE COUNT      34 a      50 c      35 g      54 t      2 others
ORIGIN
Query Match      80.8%; Score 20.2; DB 5; Length 175;
Best Local Similarity 88.0%; Pred. No. 52;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 GCTTCTTGTGAGAGCTCTGCCA 25
|||||
Db 140 GCTTCTTGTGCCAAGAGCTCTGCCA 164

Search completed: March 17, 2003, 11:31:57
Job time : 256.108 secs

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GenCore version 5.1.4\_p5\_4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 09:42:12 ; Search time 143.28 Seconds

(without alignments)  
392.938 Million cell updates/sec

Title: US-09-836-439-5

Perfect score: 25

Sequence: 1 gcttcttctgctgagagcttctca 25

Scoring table: IDENTITY-NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 112599159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	18.2	72.8	807	21 AAC48879	Arabidopsis thaliana
C 2	18.2	72.8	906	23 AAS77337	DNA encoding novel
C 3	18.2	72.8	3254	22 AAC85295	Mouse SPANK cDNA.
C 4	18.2	72.8	6912	22 AAS36854	Human cardiovascular
C 5	17.8	71.2	702	23 ABL26659	Drosophila melanog
C 6	17.8	71.2	756	21 AAH51547	Human MESTT1 relat
C 7	17.8	71.2	1000	21 AAH51581	Human cDNA encodin
C 8	17.8	71.2	1000	22 AAS30820	Human ReproSA-1 co
C 9	17.8	71.2	2573	19 AAV59153	

10	17.8	71.2	3153	23 ABL26658	Drosophila melanog
11	17.8	71.2	3870	23 ABL26632	Drosophila melanog
12	17.8	71.2	8002	24 AAD31629	Arabidopsis thaliana
C 13	17.6	70.4	223	21 AAC07949	Human secreted pro
C 14	17.6	70.4	342	24 ABL69358	Prostate cancer re
C 15	17.6	70.4	384	23 ABL11069	Human prostate exp
C 16	17.6	70.4	403	24 ABL26150	Human OREF polynuc
C 17	17.6	70.4	437	23 ABL22215	Human prostate exp
C 18	17.6	70.4	437	23 ABL41146	Human prostate exp
C 19	17.6	70.4	437	23 ABL41150	Human prostate exp
C 20	17.6	70.4	474	23 ABA92961	Human cDNA clone C
C 21	17.6	70.4	745	23 AAS41246	Human normal ovar
C 22	17.6	70.4	749	23 AAS88970	DNA encoding novel
C 23	17.6	70.4	761	20 AA242214	Human normal blad
C 24	17.6	70.4	779	21 AAC78068	Human cancer assoc
C 25	17.6	70.4	5252	22 AAF30010	Rat CARD-6 cDNA.
C 26	17.6	70.4	23899	23 ABL10362	Drosophila melanog
C 27	17.4	69.6	2849	24 ABL53927	Human dihydroorota
C 28	17.2	68.8	570	22 AAL13764	Human breast cance
C 29	17.2	68.8	687	21 AAF13683	Aspergillus oryzae
C 30	17.2	68.8	712	22 AAF22438	Human breast cance
C 31	17.2	68.8	712	22 AAF22514	Human breast cance
C 32	17.2	68.8	811	22 AAL22631	Human breast cance
C 33	17.2	68.8	850	21 AAF4643	Partial sequence M
C 34	17.2	68.8	904	21 AAF4645	Partial sequence M
C 35	17.2	68.8	1512	22 AAF30257	Human cDNA encodin
C 36	17.2	68.8	1578	22 AAF5927	Human polynucleot
C 37	17.2	68.8	1676	21 AAC75873	Human ORFX ORP1428
C 38	17.2	68.8	1739	22 AAH94496	Human full-length
C 39	17.2	68.8	1828	22 AAH15348	Human cDNA sequenc
C 40	17.2	68.8	1934	22 AAD12593	Human protein havt
C 41	17.2	68.8	1945	22 AAD4905	Human secreted pro
C 42	17.2	68.8	1945	24 ABL0687	Human polynucleot
C 43	17.2	68.8	2015	22 ABL0687	Nucleotide sequenc
C 44	17.2	68.8	2015	22 AAB91020	Human secreted pro
C 45	17.2	68.8	2015	22 AAF44267	Human PRO7170 nucl

## ALIGNMENTS

RESULT 1	
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ID AAC48879 standard; DNA: 807 BP.	
XX AAC48879;	
XX	
18-OCT-2000 (first entry)	
XX	
Arabidopsis thaliana DNA fragment SEQ ID NO: 59111.	
XX	
Hybridisation assay; genetic mapping; gene expression control;	
KW protein identification; signal transduction pathway;	
KW metabolic pathway; promoter; termination sequence; ss.	
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Arabidopsis thaliana.	
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EP1033405-A2.	
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25-FEB-2000; 2000EP-0301439.	
PF	
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25-FEB-1999; 9905-0121825.	
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PR 04-AUG-1999; 99US-0147302.  
PR 05-AUG-1999; 99US-0147192.  
PR 05-AUG-1999; 99US-0147260.  
PR 06-AUG-1999; 99US-0147303.  
PR 06-AUG-1999; 99US-0147416.  
PR 09-AUG-1999; 99US-0147493.  
PR 09-AUG-1999; 99US-0147935.  
PR 10-AUG-1999; 99US-0148171.  
PR 11-AUG-1999; 99US-0148319.  
PR 12-AUG-1999; 99US-0148341.  
PR 13-AUG-1999; 99US-0148565.  
PR 13-AUG-1999; 99US-0148684.  
PR 16-AUG-1999; 99US-0149368.  
PR 17-AUG-1999; 99US-0149175.  
PR 18-AUG-1999; 99US-0149426.  
PR 20-AUG-1999; 99US-0149722.  
PR 20-AUG-1999; 99US-0149723.  
PR 20-AUG-1999; 99US-0149929.  
PR 23-AUG-1999; 99US-0149902.  
PR 23-AUG-1999; 99US-0149930.  
PR 25-AUG-1999; 99US-0150566.  
PR 26-AUG-1999; 99US-0150884.  
PR 27-AUG-1999; 99US-0151065.  
PR 27-AUG-1999; 99US-0151066.  
PR 27-AUG-1999; 99US-0151080.  
PR 30-AUG-1999; 99US-0151303.  
PR 31-AUG-1999; 99US-0151303.  
PR 01-SEP-1999; 99US-0151930.  
PR 07-SEP-1999; 99US-0152363.  
PR 10-SEP-1999; 99US-0153070.  
PR 13-SEP-1999; 99US-0153758.  
PR 15-SEP-1999; 99US-0154018.  
PR 16-SEP-1999; 99US-0154039.  
PR 20-SEP-1999; 99US-0154779.  
PR 29-SEP-1999; 99US-0156596.  
PR 28-SEP-1999; 99US-0156458.  
PR 23-SEP-1999; 99US-0155139.  
PR 23-SEP-1999; 99US-0155486.  
PR 24-SEP-1999; 99US-0155659.  
PR 05-OCT-1999; 99US-0157753.  
PR 06-OCT-1999; 99US-0157865.  
PR 07-OCT-1999; 99US-0158029.  
PR 08-OCT-1999; 99US-0158232.  
PR 12-OCT-1999; 99US-0158369.  
PR 13-OCT-1999; 99US-0159293.  
PR 13-OCT-1999; 99US-0159294.  
PR 13-OCT-1999; 99US-0159295.  
PR 14-OCT-1999; 99US-0159329.



PR 11-JUN-1999; 99US-0138957.  
XX (WHEED ) WHITEHEAD INST BIOMEDICAL RES.  
PA (GEHO ) GEN HOSPITAL CORP.  
XX  
PI Chi N, Lodish HF;  
XX  
DR WPI, 2001-091404/10.  
DR P-PSDB; AAB47023.  
XX  
PT New insulin signalling protein SPANK, useful for reducing body mass,  
PT glucose intolerance or insulin resistance and for preventing or  
PT treating obesity-related and muscle-related diseases -  
XX  
PS Claim 3; Fig 5A; 65pp: English.  
XX  
CC This sequence represents the mouse SPANK cDNA. The SPANK  
CC protein comprises 3 domains:  
CC (a) a SAM (sterile alpha motif) domain;  
CC (b) a PAM (poly adenosine diphosphate-ribose polymerase) catalytic  
CC domain; and  
CC (c) an ANK domain composed of ankyrin repeats.  
CC SPANK is a cytosolic protein which can poly(ADP-ribosyl)ate itself.  
CC SPANK binds insulin-responsive aminopeptidase (IRAP) and modulates  
CC translocation of GLUT4 in the perinuclear region of adipocytes. It  
CC is an effector in the insulin signalling pathway in eukaryotic cells.  
CC SPANK is useful for reducing body mass, reducing glucose  
CC intolerance or insulin resistance, for preventing or treating  
CC obesity-related diseases or disorders, such as obesity, cardiac  
CC insufficiency, coronary insufficiency, stroke, hypertension,  
CC atheromatous disease, atherosclerosis, high blood pressure, non-insulin  
CC dependent diabetes, hyperlipidaemia, hyperuricaemia and Syndrome X and is  
CC also useful for preventing or treating muscle-related diseases or  
CC disorders, such as muscular dystrophy, muscle atrophy and muscle  
CC fatigue. Antibodies immunospecific for SPANK are useful for detecting  
CC the presence of SPANK polypeptide in a biological sample.  
XX  
SQ Sequence 3254 BP; 871 A; 746 C; 863 G; 774 T; 0 other;  
XX  
Query Match 72.8%; Score 18.2; DB 22; Length 3254;  
Best Local Similarity 87.0%; Pred. No. 1.7e+02;  
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
XX  
QY 3 TTCTTGTGCTGAGAGCTCTTCCA 25  
II |||||||||  
Db 2930 TTATTTGCTGAGAGCTCTTCCA 2952  
XX  
RESULT 4  
ID AAS36854 standard; DNA; 6912 BP.  
XX  
AC AAS36854;  
XX  
DT 17-DEC-2001 (first entry)  
XX  
DE Human cardiovascular system antigen genomic DNA SEQ ID No 2354.  
XX  
KW Cardiovascular system antigen; human; mouse; rabbit; goat; horse; cat;  
KW chicken; sheep; immunosuppressive; antiarthritic; vasotropic; dog;  
KW antihemorrhagic; antiproliferative; cytostatic; cardiant; neuroprotective;  
KW cerebroprotective; nootropic; antibacterial; virucide; fungicide; cancer;  
KW ophthalmological; vulnery; gene therapy; autoimmune disease; neoplasm;  
KW hyperproliferative disorder; breast; liver; cardiovascular disorder; ds;  
KW cerebrovascular disorder; nervous system disorder; bacterial infection;  
KW fungal infection; viral infection; ocular disorder; endocrine disorder;  
KW gastrointestinal disorder; renal disorder; respiratory disorder;  
KW wound healing; skin aging; organ transplantation; tissue regeneration;  
KW anti-infertility.  
XX  
OS Homo sapiens.  
XX  
PN WO200155321-A2.

XX 02-AUG-2001.  
PD 17-JAN-2001; 2001WO-US01340.  
XX  
PR 31-JAN-2000; 2000US-0179065.  
XX 04-FEB-2000; 2000US-0180628.  
PR 24-FEB-2000; 2000US-0184664.  
PR 02-MAR-2000; 2000US-0186350.  
PR 16-MAR-2000; 2000US-0189874.  
PR 17-MAR-2000; 2000US-0190076.  
PR 18-APR-2000; 2000US-0198123.  
PR 19-MAY-2000; 2000US-0205515.  
PR 07-JUN-2000; 2000US-0209467.  
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PR 11-JUL-2000; 2000US-0217487.  
PR 11-JUL-2000; 2000US-0217496.  
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PR 26-JUL-2000; 2000US-0220963.  
PR 26-JUL-2000; 2000US-0220964.  
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PR 14-AUG-2000; 2000US-0224519.  
PR 14-AUG-2000; 2000US-0225213.  
PR 14-AUG-2000; 2000US-0225214.  
PR 14-AUG-2000; 2000US-0225266.  
PR 14-AUG-2000; 2000US-0225267.  
PR 14-AUG-2000; 2000US-0225268.  
PR 14-AUG-2000; 2000US-0225270.  
PR 14-AUG-2000; 2000US-0225447.  
PR 14-AUG-2000; 2000US-0225757.  
PR 14-AUG-2000; 2000US-0225758.  
PR 14-AUG-2000; 2000US-0225759.  
PR 18-AUG-2000; 2000US-0226279.  
PR 22-AUG-2000; 2000US-0226681.  
PR 22-AUG-2000; 2000US-0226868.  
PR 22-AUG-2000; 2000US-0227182.  
PR 23-AUG-2000; 2000US-0227009.  
PR 30-AUG-2000; 2000US-0228924.  
PR 01-SEP-2000; 2000US-0229287.  
PR 01-SEP-2000; 2000US-0229343.  
PR 01-SEP-2000; 2000US-0229344.  
PR 01-SEP-2000; 2000US-0229345.  
PR 05-SEP-2000; 2000US-0229509.  
PR 05-SEP-2000; 2000US-0229513.  
PR 06-SEP-2000; 2000US-0230437.  
PR 06-SEP-2000; 2000US-0230438.  
PR 08-SEP-2000; 2000US-0231242.  
PR 08-SEP-2000; 2000US-0231243.  
PR 08-SEP-2000; 2000US-0231244.  
PR 08-SEP-2000; 2000US-0231413.  
PR 08-SEP-2000; 2000US-0231414.  
PR 08-SEP-2000; 2000US-0232080.  
PR 08-SEP-2000; 2000US-0232081.  
PR 12-SEP-2000; 2000US-0231968.  
PR 14-SEP-2000; 2000US-0232397.  
PR 14-SEP-2000; 2000US-0232398.  
PR 14-SEP-2000; 2000US-0232399.  
PR 14-SEP-2000; 2000US-0232400.  
PR 14-SEP-2000; 2000US-0232401.  
PR 14-SEP-2000; 2000US-0233063.  
PR 14-SEP-2000; 2000US-0233064.  
PR 14-SEP-2000; 2000US-0233065.  
PR 21-SEP-2000; 2000US-0234223.  
PR 21-SEP-2000; 2000US-0234274.  
PR 25-SEP-2000; 2000US-0234997.  
PR 25-SEP-2000; 2000US-0234998.  
PR 26-SEP-2000; 2000US-0235484.  
PR 27-SEP-2000; 2000US-0235834.  
PR 27-SEP-2000; 2000US-0235836.  
PR 29-SEP-2000; 2000US-0236327.

PR 29-SEP-2000; 2000US-0236367.  
 PR 29-SEP-2000; 2000US-0236368.  
 PR 29-SEP-2000; 2000US-0236369.  
 PR 29-SEP-2000; 2000US-0236370.  
 PR 02-OCT-2000; 2000US-0236882.  
 PR 02-OCT-2000; 2000US-0237037.  
 PR 02-OCT-2000; 2000US-0237038.  
 PR 02-OCT-2000; 2000US-0237039.  
 PR 02-OCT-2000; 2000US-0237040.  
 PR 13-OCT-2000; 2000US-0239935.  
 PR 13-OCT-2000; 2000US-0239937.  
 PR 20-OCT-2000; 2000US-0240960.  
 PR 20-OCT-2000; 2000US-0241221.  
 PR 20-OCT-2000; 2000US-0241785.  
 PR 20-OCT-2000; 2000US-0241786.  
 PR 20-OCT-2000; 2000US-0241787.  
 PR 20-OCT-2000; 2000US-0241808.  
 PR 20-OCT-2000; 2000US-0241809.  
 PR 01-NOV-2000; 2000US-0241826.  
 PR 08-NOV-2000; 2000US-0244617.  
 PR 08-NOV-2000; 2000US-0246474.  
 PR 08-NOV-2000; 2000US-0246475.  
 PR 08-NOV-2000; 2000US-0246476.  
 PR 08-NOV-2000; 2000US-0246477.  
 PR 08-NOV-2000; 2000US-0246478.  
 PR 08-NOV-2000; 2000US-0246523.  
 PR 08-NOV-2000; 2000US-0246524.  
 PR 08-NOV-2000; 2000US-0246525.  
 PR 08-NOV-2000; 2000US-0246526.  
 PR 08-NOV-2000; 2000US-0246527.  
 PR 08-NOV-2000; 2000US-0246528.  
 PR 08-NOV-2000; 2000US-0246532.  
 PR 08-NOV-2000; 2000US-0246609.  
 PR 08-NOV-2000; 2000US-0246610.  
 PR 08-NOV-2000; 2000US-0246611.  
 PR 08-NOV-2000; 2000US-0246613.  
 PR 17-NOV-2000; 2000US-0249207.  
 PR 17-NOV-2000; 2000US-0249208.  
 PR 17-NOV-2000; 2000US-0249209.  
 PR 17-NOV-2000; 2000US-0249210.  
 PR 17-NOV-2000; 2000US-0249211.  
 PR 17-NOV-2000; 2000US-0249212.  
 PR 17-NOV-2000; 2000US-0249213.  
 PR 17-NOV-2000; 2000US-0249214.  
 PR 17-NOV-2000; 2000US-0249215.  
 PR 17-NOV-2000; 2000US-0249216.  
 PR 17-NOV-2000; 2000US-0249217.  
 PR 17-NOV-2000; 2000US-0249218.  
 PR 17-NOV-2000; 2000US-0249244.  
 PR 17-NOV-2000; 2000US-0249245.  
 PR 17-NOV-2000; 2000US-0249264.  
 PR 17-NOV-2000; 2000US-0249265.  
 PR 17-NOV-2000; 2000US-0249297.  
 PR 17-NOV-2000; 2000US-0249299.  
 PR 01-DEC-2000; 2000US-0249300.  
 PR 01-DEC-2000; 2000US-0250160.  
 PR 05-DEC-2000; 2000US-0250391.  
 PR 05-DEC-2000; 2000US-0251030.  
 PR 05-DEC-2000; 2000US-0251988.  
 PR 05-DEC-2000; 2000US-0256719.  
 PR 06-DEC-2000; 2000US-0251479.  
 PR 08-DEC-2000; 2000US-0251856.  
 PR 08-DEC-2000; 2000US-0251868.  
 PR 08-DEC-2000; 2000US-0251869.  
 PR 08-DEC-2000; 2000US-0251989.  
 PR 11-DEC-2000; 2000US-0251990.  
 PR 05-JAN-2001; 2001US-0259678.  
 XX  
 XX  
 PA (HUMA-) HUMAN GENOME SCI INC.  
 PI Rosen Ca, Barash SC, Ruben SM.  
 XX

DR WPI; 2001-451930/48.  
 XX  
 PT New cardiovascular system related polynucleotides and polypeptides,  
 PT useful for diagnosing, treating and/or preventing disorders of the  
 PT cardiovascular system -  
 XX  
 PS Claim 1; SEQ ID NO 2354; 674pp; English.  
 XX  
 CC Sequences AAS35741-AAS36942 represent genomic DNA molecules, which encode  
 CC the cardiovascular system antigen polypeptides of the invention.  
 CC Cardiovascular system antigens and their associated polynucleotides are  
 CC useful in the diagnosis, treatment and prevention of various types of  
 CC disorders in e.g. humans, mice, rabbits, goats, horses, cats, dogs,  
 CC chickens or sheep. A pathological condition can be determined by  
 CC detecting the presence or absence of a mutation in a cardiovascular  
 CC system antigen polynucleotide. The treatable disorders include autoimmune  
 CC diseases such as rheumatoid arthritis, hyperproliferative disorders such  
 CC as neoplasms of the breast or liver, cardiovascular disorders such as  
 CC cardiac arrest, cerebrovascular disorders such as cerebral ischemia,  
 CC nervous system disorders such as Alzheimer's disease, infections caused  
 CC by bacteria, viruses and fungi, ocular disorders such as corneal  
 CC infection, endocrine disorders such as premature labour and infertility,  
 CC gastrointestinal disorders such as Crohn's disease, renal disorders such  
 CC as glomerulonephritis and respiratory disorders such as asthma and  
 CC pleurisy. The polypeptides can also be used to aid wound healing, to  
 CC prevent skin aging due to sunburn, to maintain organs before  
 CC transplantation, to regenerate tissues and in chemotaxis.  
 CC Note: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pct\_sequences.  
 XX

Query Match 72.8%; Score 18.2; DB 22; Length 6912;  
 Best Local Similarity 87.0%; Pred. No. 1.9e+02;  
 Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 Oy 3 TTCTTCTGCTGAGAGCTCTTCCA 25  
 ||||| ||||| || |||||  
 Db 647 TTCTTCTGCTGATFAGCGCTTCCA 669

RESULT 5  
 ABL26659 standard; DNA; 702 BP.  
 ID ABL26659;  
 AC ABL26659;  
 XX  
 DT 26-MAR-2002 (first entry)  
 XX  
 DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 31450.  
 XX  
 KM Drosophila: developmental biology; cell signalling; insecticide;  
 KM pharmaceutical; gene; ds.  
 XX  
 OS Drosophila melanogaster.  
 XX  
 PN WO200171042-A2.  
 PD 27-SEP-2001.  
 XX  
 PF 23-MAR-2001; 2001WO-US09231.  
 XX  
 PR 23-MAR-2000; 2000US-191637P.  
 PR 11-JUL-2000; 2000US-0614150.  
 XX  
 PA (PEKE ) PE CORP NY.  
 XX  
 PI Venter JC, Adams M, Li PWD, Myers EW;  
 XX  
 DR WPI; 2001-656860/75.  
 XX  
 PT New isolated nucleic acid detection reagent for detecting 1000 or more  
 PT genes from Drosophila and for elucidating cell signalling and cell-cell

PT Interactions -

XX

PS Claim 1; SEQ ID NO 31450; 21bp + Sequence Listing; English.

XX

CC The invention relates to an isolated nucleic acid detection reagent capable of detecting 1000 or more genes from *Drosophila*. The invention is useful in developmental biology and in elucidating cell signalling and cell-cell interactions in higher eukaryotes for the development of insecticides, therapeutics and pharmaceutical drugs. The invention discloses genomic DNA sequences (AB16176-AB130511), expressed DNA sequences (AB101840-AB16175) and the encoded proteins

CC (AB57737-AB572072).

CC The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published\_pct\_sequences.

CC

XX Sequence 702 BP; 152 A; 190 C; 158 G; 202 T; 0 other;

SQ

Query Match 71.2%; Score 17.8; DB 23; Length 702;

Best Local Similarity 90.5%; Pred. No. 2e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 TCTTTGCTGAGAGCTCTTCCA 25

||||| ||||| ||||| |||||

Db 475 TCTTTGCTGAGAGCTCTTCCA 495

RESULT 6

AAH51547/c

ID AAH51547 standard; DNA; 756 BP.

XX

AC AAH51547;

XX

DE 29-AUG-2001 (first entry)

XX

DE Human MGSTII related DNA containing a biallelic polymorphism SEQ ID 438.

XX

KW Human; biallelic marker; single nucleotide polymorphism; SNP; MGSTII; microsomal glutathione S-transferase II; malate decarboxylase enzyme; DME1; ME1; cytochrome P450; glutathione reductase; GSHR; GSHS; GGT5; flavin-containing monooxygenase; FMO; gamma-glutamyltransferase 5; dipeptidase; DP; glucose 6-phosphate dehydrogenase; G6PDH; haplotype; phosphogluconate dehydrogenase; PGDH; drug metabolism; phenotype; uridine diphosphate glucuronosyl transferase; UGT2; asthma; hepatotoxicity; zileuton; ds.

XX

OS Homo sapiens.

XX

PN WO200058508-A2.

XX

PD 05-OCT-2000.

XX

PE 24-MAR-2000; 2000WO-IB00403.

XX

PR 25-MAR-1999; 99US-0126269.

XX

PR 30-APR-1999; 99US-0131961.

XX

PA (GEST ) GENSET.

XX

PI Blumenfeld M, Bougueleret L, Chumakov I, Cohen-Akenine A;

XX

DR WPI; 2000-638353/61.

XX

PT Polynucleotides comprising sequences from malate decarboxylase enzyme-related biallelic markers used for genotyping -

XX

PS Claim 13; Page 616; 673bp; English.

XX

CC Sequences AAH51110-AAH51593 represent human DNA fragments which contain biallelic markers. The sequences are related to various human genes including microsomal glutathione S-transferase II (MGSTII), malate decarboxylase enzyme (DME1/ME1), cytochrome P450, glutathione reductase/synthase (GSHR/GSHS), flavin-containing monooxygenases (FMO),

CC gamma-glutamyltransferase 5 (GGT5), dipeptidase (DP), glucose 6-phosphate dehydrogenase (G6PDH), phosphogluconate dehydrogenase (PGDH), and uridine diphosphate glucuronosyl transferases (UGT2). Each of these sequences contains a biallelic marker/polymorphism, which is represented in the sequence as a degenerate/undefined base. The genes to which the biallelic marker containing sequences are related are involved in drug metabolism.

CC Sequences AAH51594 - AAH51598 represent the genomic sequence of the MGSTII gene and four alternative MGSTII cDNA sequences. AAB62905-AAH62906 are MGSTII gene products. PCR primers AAH51599 and AAH51600 are used in an example for the amplification of human genomic DNA fragments. The invention includes a method of genotyping comprising determining the identity of a nucleotide at a DME- or MGSTII-related biallelic marker in a biological sample. The method is used to determine the frequency in population of an allele of a DME- or MGST-II related biallelic marker and to select an individual for inclusion in a clinical trial of a drug treatment. The method is also used to detect association between allele CC and phenotype, and to detect association between haplotype and phenotype. The polynucleotides are used, in hybridization assays, sequencing assays CC or allele specific amplification assays. The method can be used to determine whether an individual suffers or is at risk of developing CC asthma or is at risk of developing hepatotoxicity on treatment with CC zileuton.

XX

SQ Sequence 756 BP; 224 A; 140 C; 177 G; 213 T; 2 other;

XX

Query Match 71.2%; Score 17.8; DB 21; Length 756;

Best Local Similarity 90.5%; Pred. No. 2e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TTTCCTTGCTGAGAGCTCTTC 23

||||| ||||| ||||| |||||

Db 614 TTTCCTTGCTGAGAGCTCTTC 594

RESULT 7

AAH51581/c

ID AAH51581 standard; DNA; 1000 BP.

XX

AC AAH51581;

XX

DE 29-AUG-2001 (first entry)

XX

DE Human MGSTII related DNA containing a biallelic polymorphism SEQ ID 472.

XX

KW Human; biallelic marker; single nucleotide polymorphism; SNP; MGSTII; microsomal glutathione S-transferase II; malate decarboxylase enzyme; DME1; ME1; cytochrome P450; glutathione reductase; GSHR; GSHS; GGT5; flavin-containing monooxygenase; FMO; gamma-glutamyltransferase 5; dipeptidase; DP; glucose 6-phosphate dehydrogenase; G6PDH; haplotype; phosphogluconate dehydrogenase; PGDH; drug metabolism; phenotype; uridine diphosphate glucuronosyl transferase; UGT2; asthma; hepatotoxicity; zileuton; ds.

XX

OS Homo sapiens.

XX

PN WO200058508-A2.

XX

PD 05-OCT-2000.

XX

PE 24-MAR-2000; 2000WO-IB00403.

XX

PR 25-MAR-1999; 99US-0126269.

XX

PR 30-APR-1999; 99US-0131961.

XX

PA (GEST ) GENSET.

XX

PI Blumenfeld M, Bougueleret L, Chumakov I, Cohen-Akenine A;

XX

DR WPI; 2000-638353/61.

XX

PT Polynucleotides comprising sequences from malate decarboxylase enzyme-related biallelic markers used for genotyping -

XX

PS Claim 13; Page 641; 673pp; English.

XX Sequences AAH51110-AAH51593 represent human DNA fragments which contain

CC biallelic markers. The sequences are related to various human genes

CC including microsomal glutathione S-transferase II (MGSTII), malate

CC decarboxylase enzyme (DME/ME1), cytochrome P450, glutathione

CC reductase/synthase (GSHR/GSHS), flavin-containing monooxygenases (FMO),

CC gamma-glutamyltransferase 5 (GGT5), dipeptidase (DP), glucose 6-phosphate

CC dehydrogenase (G6PDH), phosphogluconate dehydrogenase (PGDH), and uridine

CC diphosphate glucuronosyl transferases (UGT2). Each of these sequences

CC contains a biallelic marker/polymorphism, which is represented in the

CC sequence as a degenerate/undefined base. The genes to which the biallelic

CC marker containing sequences are related are involved in drug metabolism.

CC Sequences AAH51594 - AAH51598 represent the genomic sequence of the

CC MGSTII gene and four alternative MGSTII cDNA sequences. AAB62905-AAB62906

CC are MGSTII gene products. PCR primers AAH51599 and AAH51600 are used in

CC an example for the amplification of human genomic DNA fragments. The

CC invention includes a method of genotyping comprising determining the

CC identity of a nucleotide at a DME- or MGSTII-related biallelic marker in

CC a biological sample. The method is used to determine the frequency in

CC a population of an allele of a DME- or MGSTII-related biallelic marker and

CC to select an individual for inclusion in a clinical trial of a drug

CC treatment. The method is also used to detect association between allele

CC and phenotype, and to detect association between haplotype and phenotype.

CC The polynucleotides are used, in hybridization assays, sequencing assays

CC or allele specific amplification assays. The method can be used to

CC determine whether an individual suffers or is at risk of developing

CC asthma or is at risk of developing hepatotoxicity on treatment with

CC zileuton.

XX

SQ Sequence 1000 BP; 300 A; 200 C; 211 G; 288 T; 1 other;

Query Match 71.2%; Score 17.8; DB 21; Length 1000;

Best Local Similarity 90.5%; Pred. No. 2.1e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 TTCTTTGCTGAGAGCTCTTC 23

DB 851 TTCTTTGCTGAGAGCTCTTC 831

RESULT 8

AA530820/c

ID AA530820 standard; cDNA; 1000 BP.

XX

AC AA530820;

XX

DT 04-DEC-2001 (first entry)

XX

DE Human cDNA encoding G protein-coupled receptor ngPCR-2428.

XX

XX Human; G protein-coupled receptor; ngPCR-x; ss; antiviral; analgesic;

KW cytototoxic; cardiatic; antidiabetic; anorectic; hypotensive; hypertensive;

KW antiparkinsonian; nootropic; neuroprotective; antidepressant;

KW viral infection; HIV-1; human immunodeficiency virus; HIV-2; pain;

KW cancer; metabolic disease; cardiovascular disease; type 2 diabetes;

KW obesity; anorexia; hypotension; hypertension; myocardial infarction;

KW atherosclerosis; Parkinson's disease; psychosis; neurological disorder;

KW schizophrenia; migraine; major depression; anxiety; mental disorder;

XX manic depression; dyskinesia; Huntington's disease; Tourette's Syndrome.

XX

OS Homo sapiens.

XX

PN WO200166750-A2.

XX

PD 13-SEP-2001.

XX

PF 08-MAR-2001; 2001WO-US07322.

XX

PR 08-MAR-2000; 2000US-0187581.

PR 08-MAR-2000; 2000US-0187582.

PR 08-MAR-2000; 2000US-0187714.

PR 08-MAR-2000; 2000US-0187715.

PR 08-MAR-2000; 2000US-0187825.

PR 08-MAR-2000; 2000US-0187828.

PR 08-MAR-2000; 2000US-0187829.

PR 08-MAR-2000; 2000US-0187830.

PR 08-MAR-2000; 2000US-0187833.

PR 08-MAR-2000; 2000US-0187874.

PR 08-MAR-2000; 2000US-0187930.

PR 08-MAR-2000; 2000US-0188049.

PR 08-MAR-2000; 2000US-0189294.

PR 08-MAR-2000; 2000US-0189299.

PR 08-MAR-2000; 2000US-0187928.

XX

PA (PMAA ) PHARMACIA & UPJOHN CO.

XX

PL Vogel1 G, Wood LS;

XX

DR WPI: 2001-536778/59.

DR P-PSDB; AAU19251.

XX

PT Isolated nucleic acid molecules encoding G protein-coupled receptors

PT termed ngPCR-x, useful in the treatment and diagnosis of viral

PT infections, cancers and mental disorders (e.g. Parkinson's disease and

PT schizophrenia) -

XX

PS Claim 4; Page 221-222; 336pp; English.

XX

CC The invention relates to novel isolated nucleic acid molecules encoding

CC G protein-coupled receptors termed ngPCR-x, ngPCR-x polynucleotides,

CC polypeptides, and modulators may be used in the treatment of diseases and

CC conditions such as infections, such as viral infections caused by HIV-1

CC (human immunodeficiency virus) or HIV-2, pain, cancers, metabolic and

CC cardiovascular diseases and disorders (e.g., type 2 diabetes, obesity,

CC anorexia, hypotension, hypertension, myocardial infarction,

CC atherosclerosis), Parkinson's disease, and psychotic and

CC neurological disorders, including schizophrenia, migraine, major

CC depression, anxiety, mental disorder, manic depression, and

CC dyskinesias, such as Huntington's disease or Tourette's Syndrome

CC and many other diseases and syndromes listed in the specification.

CC ngPCR-x polynucleotides and polypeptides, as well as ngPCR-x

CC modulators, may also be used in diagnostic assays for such diseases or

CC conditions. The present sequence encodes a G protein-coupled

CC receptor of the invention.

XX

SQ Sequence 1000 BP; 265 A; 208 C; 299 G; 228 T; 0 other;

Query Match 71.2%; Score 17.8; DB 22; Length 1000;

Best Local Similarity 90.5%; Pred. No. 2.1e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 TTCTTTGCTGAGAGCTCTTC 23

DB 791 TTCTTTGCTGAGAGCTCTTC 771

RESULT 9

AAV59153

ID AAV59153 standard; cDNA; 2573 BP.

XX

AC AAV59153;

XX

DT 14-DEC-1998 (first entry)

XX

DE Human ReproSA-1 coding sequence.

XX

XX ss; human; ReproSA-1; contraceptive; anti-sperm; fertility;

KW in vitro fertilisation.

XX

XX Homo sapiens.

OS

XX

XX Key Location/Qualifiers

XX CDS 125..1654

FT /\*tag= a

FT /\*product= "ReproSA-1 protein"

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XX XX WO9836073-A1.
XX XX 20-AUG-1998.
XX PF 17-FEB-1998; 98WO-US03513.
XX PR 18-FEB-1997; 97US-0039574.
XX PA (REPR-) REPROGEN INC.
XX PI French CK, Neilson LI, Schneider PA, Yamamoto KK;
XX DR WPI; 1998-480771/41.
XX DR P-PSDB; AAM77323.
XX PT Sperm-specific polypeptide ReproSA-1 and related nucleic acid -
XX PT transformed cells and antibodies, useful in contraceptive vaccines
XX PT or compositions, and for diagnosis and treatment of infertility
XX PS Claim 16; Fig 1; 62pp; English.
XX CC The ReproSA-1 polypeptide, its fragments, or nucleic acid encoding them
XX CC are used in contraceptive vaccines to generate an anti-sperm response.
XX CC Antibodies can also be used as topical contraceptive (spermicide), also
XX CC antisense sequences, ribozymes and triplex-forming molecules. Detection,
XX CC by reaction with immobilised peptides, of anti-(I) antibodies (in serum,
XX CC semen, saliva, cervical or vaginal mucosa) is used to diagnose
XX CC immunological infertility and to monitor the effect of vaccination.
XX CC nucleotide fragments are used to diagnose infertility-associated
XX CC mutations in the gene encoding ReproSA, and for isolation of related
XX CC sequences. Peptides and anti-idiotypic antibodies against the
XX CC antibodies are used to inactivate the antibodies i.e. to increase
XX CC fertility and may be added to sperm samples before in vitro
XX CC fertilisation.
XX SO Sequence 2573 BP; 754 A; 532 C; 586 G; 701 T; 0 other;

Query Match 71.2%; Score 17.8; DB 19; Length 2573;
Best Local Similarity 90.5%; Pred. No. 2.5e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TTCTTTGCTGAGAGCTCTTC 23
DB 252 TGTCTTGTGAGACTCTTC 272

RESULT 10
ABL26658
ID ABL26658 standard; DNA; 3153 BP.
XX AC ABL26658;
XX XX 26-MAR-2002 (first entry)
XX DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 31447.
XX XX Drosophila; developmental biology; cell signalling; insecticide;
XX KM pharmaceutical; gene; ds.
XX XX Drosophila melanogaster.
XX OS WO200171042-A2.
XX PN 27-SEP-2001.
XX PD 23-MAR-2001; 2001WO-US09231.
XX PF 23-MAR-2000; 2000US-191637P.
XX PR 11-JUL-2000; 2000US-0614150.
XX PA (PEKE ) PE CORP NY.
XX XX

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PI Venter JC, Adams M, Li PWD, Myers EW;
XX XX WPI; 2001-656860/75.
XX DR New isolated nucleic acid detection reagent for detecting 1000 or more
XX PT genes from Drosophila and for elucidating cell signalling and cell-cell
XX PT interactions -
XX PS Claim 1; SEQ ID NO 31447; 21pp + Sequence Listing; English.
XX CC The invention relates to an isolated nucleic acid detection reagent
XX CC capable of detecting 1000 or more genes from Drosophila. The invention is
XX CC useful in developmental biology and in elucidating cell signalling and
XX CC cell-cell interactions in higher eukaryotes for the development of
XX CC insecticides, therapeutics and pharmaceutical drugs. The invention
XX CC discloses genomic DNA sequences (AB116176-AB130511), expressed DNA
XX CC sequences (AB57737-AB572072).
XX CC The sequence data for this patent did not form part of the printed
XX CC specification, but was obtained in electronic format directly from WIPO
XX CC at ftp.wipo.int/pub/published_pct_sequences.
XX SO Sequence 3153 BP; 864 A; 693 C; 665 G; 931 T; 0 other;

Query Match 71.2%; Score 17.8; DB 23; Length 3153;
Best Local Similarity 90.5%; Pred. No. 2.6e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 TCTTTGCTGAGAGCTCTTCCA 25
DB 1746 TCTTCTGTGAGCGCTTCCA 1766

RESULT 11
ABL26632
ID ABL26632 standard; DNA; 3870 BP.
XX AC ABL26632;
XX XX 26-MAR-2002 (first entry)
XX DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 31369.
XX XX Drosophila; developmental biology; cell signalling; insecticide;
XX KM pharmaceutical; gene; ds.
XX XX Drosophila melanogaster.
XX OS WO200171042-A2.
XX PN 27-SEP-2001.
XX PD 23-MAR-2001; 2001WO-US09231.
XX PF 23-MAR-2000; 2000US-191637P.
XX PR 11-JUL-2000; 2000US-0614150.
XX PA (PEKE ) PE CORP NY.
XX PI Venter JC, Adams M, Li PWD, Myers EW;
XX XX WPI; 2001-656860/75.
XX DR New isolated nucleic acid detection reagent for detecting 1000 or more
XX PT genes from Drosophila and for elucidating cell signalling and cell-cell
XX PT interactions -
XX PS Claim 1; SEQ ID NO 31369; 21pp + Sequence Listing; English.
XX CC The invention relates to an isolated nucleic acid detection reagent
XX CC capable of detecting 1000 or more genes from Drosophila. The invention is
XX CC useful in developmental biology and in elucidating cell signalling and
XX CC cell-cell interactions in higher eukaryotes for the development of

```



CC Insecticides, therapeutics and pharmaceutical drugs. The invention  
CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA  
CC sequences (ABL1840-ABL16175) and the encoded proteins  
CC (ABB57737-ABB72072).  
CC The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pcl\_sequences.  
XX  
SQ Sequence 3870 BP; 961 A; 917 C; 886 G; 1106 T; 0 other;  
XX  
Query Match 71.2%; Score 17.8; DB 23; Length 3870;  
Best Local Similarity 90.5%; Pred. No. 2.7e+02;  
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
OY 5 TCTTTGCTGAGAGCTCTTCCA 25  
DB 103 TCTTCTGCTGAGAGCTCTTCCA 123  
XX  
RESULT 12  
ID AAD31629 standard; DNA; 8002 BP.  
XX  
AC AAD31629;  
XX  
DT 18-JUN-2002 (first entry)  
XX  
DE Arabidopsis thaliana GT1209 gene.  
XX  
KM GT1209 gene; herbicide; plant tissue; cell; seedling growth; development;  
XX db.  
XX Arabidopsis thaliana.  
XX  
PN W0200212273-A2.  
XX  
PD 14-FEB-2002.  
XX  
PF 01-AUG-2001; 2001WO-EP08910.  
XX  
PR 03-AUG-2000; 2000US-222779P.  
XX  
PA (SYGN ) SYNGENTA PARTICIPATIONS AG.  
XX  
PI Levin JZ, Megrich Glover L, Budziszewski GJ;  
XX  
DR WPI; 2002-241730/29.  
XX  
PT New polypeptide having GT1209, GT1354 or GT0946 activity, obtained from  
PT Arabidopsis, useful as herbicide targets in screening assays to  
PT identify the inhibitors or potential herbicides  
XX  
PS Disclosure: Page 72-74; 82pp; English.  
XX  
CC The invention relates to a polypeptide having GT1209, GT1354 or GT0946  
CC activity obtained from Arabidopsis. The invention may also be applied to  
CC the development of herbicide tolerant plants, plant tissues, plant seeds  
CC and plant cells. The polypeptide is useful as herbicide targets in  
CC screening assays to identify potential herbicides and inhibitors of  
CC GT1209, GT1354 or GT0946 activity. A compound having herbicidal activity  
CC is useful for suppressing the growth of a plant. The newly discovered  
CC GT1209, GT1354 or GT0946 genes are essential for seedling growth and  
CC development. The present sequence is Arabidopsis thaliana GT1209 gene.  
XX  
SQ Sequence 8002 BP; 2119 A; 1489 C; 1502 G; 2892 T; 0 other;  
XX  
Query Match 71.2%; Score 17.8; DB 24; Length 8002;  
Best Local Similarity 90.5%; Pred. No. 3e+02;  
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
OY 2 CTTTCTTGTGCTGAGAGCTCTT 22  
DB 6994 CTTTCTTGTGCTGATTCCTCT 7014

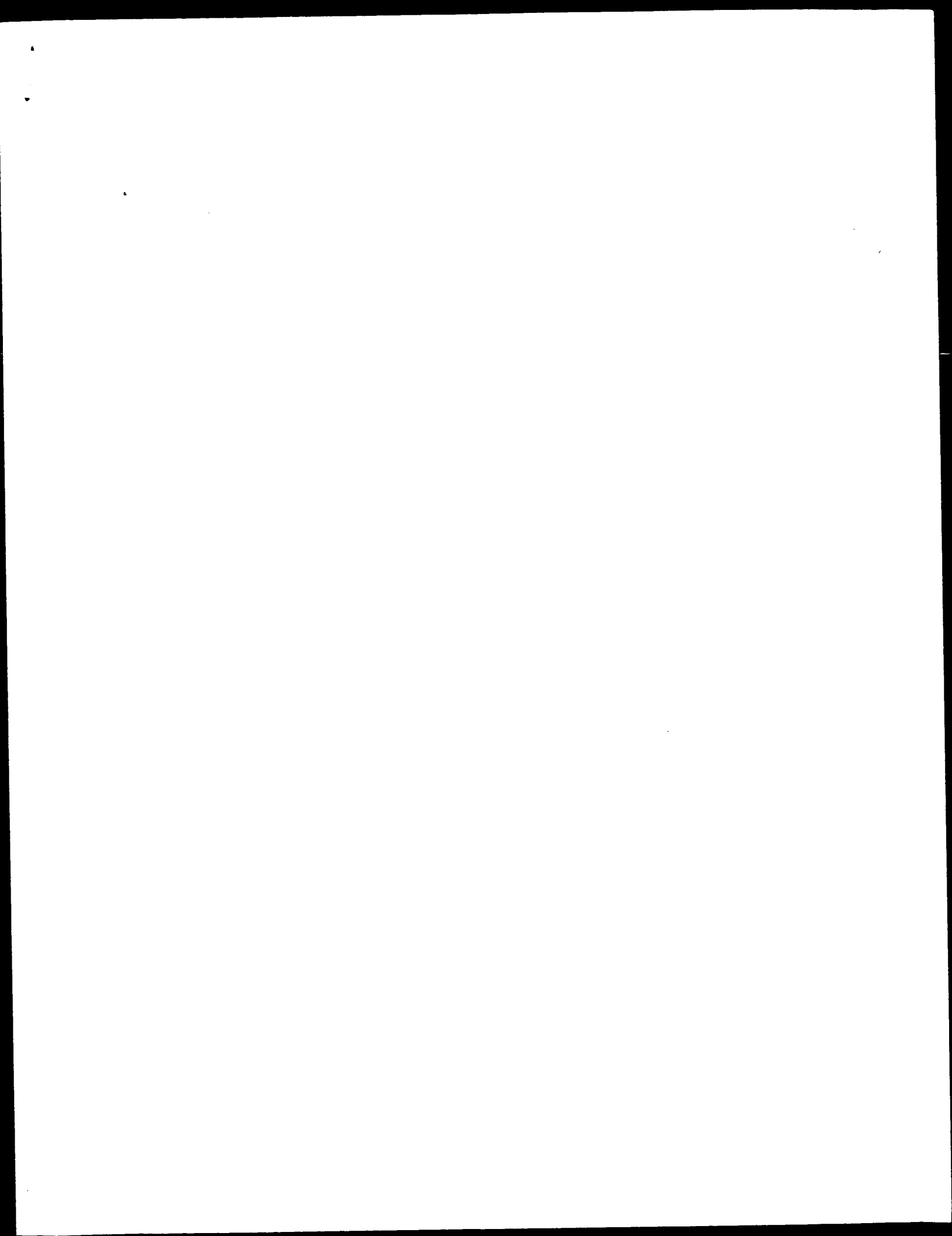
XX  
RESULT 13  
ID AAC07949/GC  
XX AAC07949 standard; cDNA; 223 BP.  
XX  
AC AAC07949;  
XX  
DT 06-OCT-2000 (first entry)  
XX  
DE Human secreted protein 5' EST, SEQ ID NO: 12024.  
XX  
KM Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;  
KM gene therapy; chromosome mapping; ss.  
XX  
OS Homo sapiens.  
XX  
PN EP1033401-A2.  
XX  
PD 06-SEP-2000.  
XX  
PF 21-FEB-2000; 2000EP-0200610.  
XX  
PR 26-FEB-1999; 99US-0122487.  
XX  
PA (GEST ) GENSET.  
XX  
PI Dumas Milne Edwards J, Duclert A, Giordano J;  
XX  
DR WPI; 2000-500381/45.  
XX  
PT New nucleic acid that is a 5' expressed sequence tag (5' EST) for  
PT obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for  
PT diagnostic, forensic, gene therapy and chromosome mapping procedures -  
XX  
PS Claim 1; SEQ ID 12024; 71pp + CD-ROM; English.  
XX  
CC The present sequence is one of a large number of 5' ESTs derived from  
CC mRNAs encoding secreted proteins. No ORF has yet been conclusively  
CC identified within the present sequence. The 5' ESTs were prepared from  
CC total human RNAs or polyA+ RNAs derived from 30 different tissues. EST  
CC sequences usually correspond mainly to the 3' untranslated region (UTR)  
CC of the mRNA because they are often obtained from oligo-dT primed cDNA  
CC libraries. Such ESTs are not well suited for isolating cDNA sequences  
CC derived from the 5' ends of mRNAs and even in those cases where longer  
CC cDNA sequences have been obtained, the full 5' UTR is rarely included.  
CC 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be  
CC used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used  
CC in diagnostic, forensic, gene therapy and chromosome mapping procedures.  
CC They are used to obtain upstream regulatory sequences and to design  
CC expression and secretion vectors.  
XX  
SQ Sequence 223 BP; 77 A; 38 C; 53 G; 54 T; 1 other;  
XX  
Query Match 70.4%; Score 17.6; DB 21; Length 223;  
Best Local Similarity 83.3%; Pred. No. 2e+02;  
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
OY 1 GCTTCTTGTGCTGAGAGCTCTTCC 24  
DB 152 GCCTTCTTGTGAGAGATTTCTTCC 129  
XX  
RESULT 14  
ID ABL69358  
XX ABL69358 standard; DNA; 342 BP.  
XX  
AC ABL69358;  
XX  
DT 15-MAY-2002 (first entry)  
XX  
DE Prostate cancer related gene sequence SEQ ID NO: 7695.  
XX

KW Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;  
 KW stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;  
 KW cytostatic; gene therapy; antineoplastic; Wilm's tumour; adenocarcinoma;  
 KW gene; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 XX WO200194629-A2.  
 XX  
 PD 13-DEC-2001.  
 XX  
 PF 30-MAY-2001; 2001WO-US10838.  
 XX  
 XX 05-JUN-2000; 2000US-209473P.  
 PR 05-JUN-2000; 2000US-209531P.  
 PR 18-SEP-2000; 2000US-233133P.  
 PR 18-SEP-2000; 2000US-233617P.  
 PR 20-SEP-2000; 2000US-234009P.  
 PR 20-SEP-2000; 2000US-234034P.  
 PR 20-SEP-2000; 2000US-234052P.  
 PR 22-SEP-2000; 2000US-234509P.  
 PR 22-SEP-2000; 2000US-234523P.  
 PR 25-SEP-2000; 2000US-234923P.  
 PR 25-SEP-2000; 2000US-234924P.  
 PR 25-SEP-2000; 2000US-235077P.  
 PR 25-SEP-2000; 2000US-235082P.  
 PR 25-SEP-2000; 2000US-235134P.  
 PR 25-SEP-2000; 2000US-235280P.  
 PR 26-SEP-2000; 2000US-235637P.  
 PR 26-SEP-2000; 2000US-235638P.  
 PR 27-SEP-2000; 2000US-235711P.  
 PR 27-SEP-2000; 2000US-235720P.  
 PR 27-SEP-2000; 2000US-235840P.  
 PR 27-SEP-2000; 2000US-235863P.  
 PR 28-SEP-2000; 2000US-236028P.  
 PR 28-SEP-2000; 2000US-236032P.  
 PR 28-SEP-2000; 2000US-236033P.  
 PR 28-SEP-2000; 2000US-236034P.  
 PR 28-SEP-2000; 2000US-236109P.  
 PR 28-SEP-2000; 2000US-236111P.  
 PR 29-SEP-2000; 2000US-236842P.  
 PR 29-SEP-2000; 2000US-236891P.  
 PR 02-OCT-2000; 2000US-237172P.  
 PR 02-OCT-2000; 2000US-237173P.  
 PR 02-OCT-2000; 2000US-237278P.  
 PR 02-OCT-2000; 2000US-237294P.  
 PR 02-OCT-2000; 2000US-237295P.  
 PR 02-OCT-2000; 2000US-237316P.  
 PR 03-OCT-2000; 2000US-237425P.  
 PR 03-OCT-2000; 2000US-237598P.  
 PR 03-OCT-2000; 2000US-237604P.  
 PR 03-OCT-2000; 2000US-237606P.  
 PR 03-OCT-2000; 2000US-237608P.  
 PR 01-NOV-2000; 2000US-244867P.  
 PR 01-NOV-2000; 2000US-245084P.  
 XX  
 PA (AVALON PHARM.  
 XX  
 PI Young PE, Augustus M, Carter KC, Edner R, Endress G, Horrigan S;  
 PI Soppet DR, Weaver Z;  
 XX  
 XX WPI; 2002-188264/24.  
 XX  
 PT Screening for anti-neoplastic agent involves exposing cells to a  
 PT chemical agent to be tested for anti-neoplastic activity, and  
 PT determining a change in expression of a gene of a signature gene set -  
 XX  
 XX Claim 1; SEQ ID 7695; 44pp; English.  
 XX  
 CC The present invention describes a method (M1) for screening for an  
 CC anti-neoplastic agent. The method involves exposing cells to a chemical  
 CC agent to be tested for anti-neoplastic activity, determining a change in  
 CC expression of at least one gene (I) of a signature gene set, where (I)

CC comprises a sequence (S) selected from 8447 sequences (given in AB161664  
 CC to AB170110), or is at least 95% identical to (S), where a change in  
 CC expression is indicative of anti-neoplastic activity. (I) has cytostatic  
 CC activity and can be used in gene therapy. M1 can be used for screening  
 CC an anti-neoplastic agent, and can be used for producing a product which  
 CC is the data collected with respect to the anti-neoplastic agent as a  
 CC result of M1, and the data is sufficient to convey the chemical  
 CC structure and/or properties of the agent. M1 can be used in the  
 CC treatment of cancer such as colon, breast, stomach, lung, thyroid,  
 CC oesophageal, ovarian, kidney, prostate or pancreatic cancer,  
 CC adenocarcinoma, carcinoma, clear cell cancer, infiltrating ductal cancer,  
 CC infiltrating lobular cancer, squamous cell carcinoma, neuroendocrine  
 CC carcinoma, papillary carcinoma and Wilm's tumour.  
 XX  
 SQ Sequence 342 BP; 36 A; 109 C; 92 G; 105 T; 0 other;  
 Query Match 70.4%; Score 17.6; DB 24; Length 342;  
 Best Local Similarity 83.3%; Pred. No. 2.2e+02;  
 Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 Qy 1 GCTTCTTGTGCTGAGAGCTCTCC 24  
 Db 67 GCTTCTTGTGCTGAGAGCTCTCC 90  
 RESULT 15  
 ID ABV1069 standard; cDNA; 384 BP.  
 AC ABV1069;  
 XX 13-SEP-2002 (first entry)  
 DT  
 XX Human prostate expression marker cDNA 11060.  
 DE  
 XX Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
 KW pharmacogenomic marker; gene; ss.  
 OS Homo sapiens.  
 XX WO200160860-A2.  
 XX  
 XX 23-AUG-2001.  
 PD  
 XX 20-FEB-2001; 2001WO-US05171.  
 PF  
 XX 17-FEB-2000; 2000US-183319P.  
 PR 16-MAR-2000; 2000US-189862P.  
 PR 25-MAY-2000; 2000US-207454P.  
 PR 09-JUN-2000; 2000US-211314P.  
 PR 18-JUL-2000; 2000US-219007P.  
 PR 13-DEC-2000; 2000US-255281P.  
 XX  
 PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
 XX  
 PI Schlegel R, Endege WO, Monahan JE;  
 PI  
 XX WPI; 2001-662795/76.  
 DR  
 XX  
 PT Novel isolated nucleic acid molecule associated with cancerous state of  
 PT prostate cells and correlating with presence of prostate cancer, useful  
 PT for detecting presence of prostate cancer, stage of prostate cancer -  
 XX  
 XX Claim 1; Page 1792; 11750pp; English.  
 PS  
 CC The invention relates to an isolated nucleic acid molecule (I) comprising  
 CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
 CC specification or its complement. (I) is useful for:  
 CC (a) assessing whether a patient is afflicted with prostate cancer;  
 CC (b) monitoring the progression of prostate cancer in a patient;  
 CC (c) assessing the efficacy of a test compound to inhibit prostate  
 CC cancer in a patient;  
 CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer

CC in a patient;  
CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
CC (f) assessing the prostate cell carcinogenic potential of a compound;  
CC (g) determining whether prostate cancer has metastasized in a patient;  
CC (h) assessing the aggressiveness or indolence of prostate cancer in a  
CC patient;  
CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.  
XX  
SQ Sequence 384 BP; 139 A; 87 C; 73 G; 85 T; 0 other;  
Query Match .70.4%; Score 17.6; DB 23; Length 384;  
Best Local Similarity 83.3%; Pred. No. 2.2e+02;  
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
Qy 2 CTTCTTTCGAGAGCTCTTCA 25  
||||||| ||||| |||  
Db 374 CTTCTTTCGAGAGCTCTTCA 351

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Job time : 146.446 secs



GenCore version 5.1.4.p5.4578  
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OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 10:23:12 ; Search time 883.737 Seconds

(without alignments)  
458.154 Million cell updates/sec

Title: US-09-836-439-5

Perfect score: 25

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Gapop 10.0, Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 08  
Maximum Match 100%

Listing first 45 summaries

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2: em\_esthum:\*  
3: em\_estlin:\*  
4: em\_estlin:\*  
5: em\_estlin:\*  
6: em\_estlin:\*  
7: em\_estlin:\*  
8: em\_estlin:\*  
9: em\_estlin:\*  
10: em\_estlin:\*  
11: em\_estlin:\*  
12: em\_estlin:\*  
13: em\_estlin:\*  
14: em\_estlin:\*  
15: em\_estlin:\*  
16: em\_estlin:\*  
17: em\_estlin:\*  
18: em\_estlin:\*  
19: em\_estlin:\*  
20: em\_estlin:\*  
21: em\_estlin:\*  
22: em\_estlin:\*  
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25: em\_estlin:\*  
26: em\_estlin:\*  
27: em\_estlin:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	23.4	93.6	402	12	BE950666 UT-M-CEO-
2	23.4	93.6	495	13	BE950666 UT-M-CEO-
3	23.4	93.6	628	13	BE950666 UT-M-CEO-
4	23.4	93.6	636	12	BE950666 UT-M-CEO-
5	23.4	93.6	649	12	BE950666 UT-M-CEO-
6	23.4	93.6	651	13	BE950666 UT-M-CEO-

7	23.4	93.6	656	13	BE950666 UT-M-CEO-
8	23.4	93.6	663	13	BE950666 UT-M-CEO-
9	23.4	93.6	672	12	BE950666 UT-M-CEO-
10	23.4	93.6	676	13	BE950666 UT-M-CEO-
11	23.4	93.6	681	13	BE950666 UT-M-CEO-
12	23.4	93.6	682	13	BE950666 UT-M-CEO-
13	23.4	93.6	684	12	BE950666 UT-M-CEO-
14	23.4	93.6	686	12	BE950666 UT-M-CEO-
15	23.4	93.6	686	12	BE950666 UT-M-CEO-
16	23.4	93.6	697	14	BE950666 UT-M-CEO-
17	23.4	93.6	717	13	BE950666 UT-M-CEO-
18	23.4	93.6	718	13	BE950666 UT-M-CEO-
19	23.4	93.6	718	13	BE950666 UT-M-CEO-
20	23.4	93.6	720	12	BE950666 UT-M-CEO-
21	23.4	93.6	743	12	BE950666 UT-M-CEO-
22	23.4	93.6	744	12	BE950666 UT-M-CEO-
23	23.4	93.6	746	12	BE950666 UT-M-CEO-
24	23.4	93.6	759	12	BE950666 UT-M-CEO-
25	23.4	93.6	761	12	BE950666 UT-M-CEO-
26	23.4	93.6	773	13	BE950666 UT-M-CEO-
27	23.4	93.6	774	12	BE950666 UT-M-CEO-
28	23.4	93.6	783	13	BE950666 UT-M-CEO-
29	23.4	93.6	790	13	BE950666 UT-M-CEO-
30	23.4	93.6	802	13	BE950666 UT-M-CEO-
31	23.4	93.6	806	12	BE950666 UT-M-CEO-
32	23.4	93.6	825	13	BE950666 UT-M-CEO-
33	23.4	93.6	832	13	BE950666 UT-M-CEO-
34	23.4	93.6	837	12	BE950666 UT-M-CEO-
35	23.4	93.6	851	13	BE950666 UT-M-CEO-
36	23.4	93.6	854	13	BE950666 UT-M-CEO-
37	23.4	93.6	873	13	BE950666 UT-M-CEO-
38	23.4	93.6	879	13	BE950666 UT-M-CEO-
39	23.4	93.6	880	14	BE950666 UT-M-CEO-
40	23.4	93.6	903	12	BE950666 UT-M-CEO-
41	23.4	93.6	909	14	BE950666 UT-M-CEO-
42	23.4	93.6	914	14	BE950666 UT-M-CEO-
43	23.4	93.6	915	13	BE950666 UT-M-CEO-
44	23.4	93.6	917	12	BE950666 UT-M-CEO-
45	23.4	93.6	918	13	BE950666 UT-M-CEO-

## ALIGNMENTS

RESULT 1  
LOCUS BE950666/c  
DEFINITION UI-M-CEO-aza-d-06-0-UI-s1 NIH BMP Ret3 Mus musculus cDNA clone  
ACCESSION BE950666  
VERSION BE950666.1 GI:10589332  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
REFERENCE 1 (bases 1 to 402)  
AUTHORS Bonaldo, M.F., Lennon, G. and Soares, M.B.  
TITLE Normalization and subtraction: two approaches to facilitate gene discovery  
JOURNAL Genome Res. 6 (9), 791-806 (1996)  
MEDLINE 97044477  
COMMENT Contact: Chin, H  
National Institute of Mental Health  
6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD  
20892-9643, USA  
Tel: 301 443 1706  
Fax: 301 443 9890  
Email: mestr@nhi.nih.gov  
The sequence contained an oligo-dT track that was present in the strand cDNA and therefore this may represent a bonafide poly A tail. The sequence tag present in the cDNA between the NciI site

and the oligo-dT track served to verify it as a clone from the retina tissue cDNA library. Preparation: M.B. Soares Lab Clone Distribution: Researchers may obtain BMAP cDNA clones from RESEARCH GENETICS. It should be noted that Bento Soares is generating a small number of additional specialized non-redundant arrays of BMAP cDNAs whose availability will be considered under appropriate and limited collaborative arrangements. The tissue for this library was contributed by Dr. Xin-Yuan Fu, Yale University School of Medicine. Seq primer: M13 Forward

# FEATURES

## Source

Location/Qualifiers  
1. 402  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UI-M-CE0-aza-d-06-0-UI"  
/clone\_lib="NIH\_BMAP\_Ret3"  
/dev\_stage="6 weeks"  
/lab\_host="DH10B (Life Technologies)"  
/note="Vector: pRT3D-Pac (Pharmacia) with a modified polylinker; Site\_1: Not I; Site\_2: Eco RI; The NIH\_BMAP\_Ret3 library is derived from mouse retina tissue. For a detailed description of the library from which this clone was derived, please visit our web site at [brunest.eng.utoronto.edu](http://brunest.eng.utoronto.edu). The tissue for this library was contributed by Dr. Xin-Yuan Fu, Yale University School of Medicine.  
TAG\_Lib=NIH\_BMAP\_Ret3  
TAG\_Tissue=adult-retina  
TAG\_SEQ=GTCCAGCGCCGAC

BASE COUNT 90 a 86 c 126 g 100 t

Query Match 93.6%; Score 23.4; DB 12; Length 402;  
Best Local Similarity 96.0%; Pred. No. 19;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTGTGAGAGCTCTTCCA 25  
|||||  
Db 334 GCTTCTTGTGAGAGCTCTTCCA 310

RESULT 2 495 bp mRNA linear EST 20-SEP-2001  
LOCUS B1737699 603358627F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:5365638 5',  
DEFINITION mRNA sequence.  
ACCESSION B1737699  
VERSION B1737699.1 GI:15714712  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Mammalia; Chordata; Craniata; Vertebrata; Euteleostomi; Eukaryota; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.  
REFERENCE NIH-MGC <http://mgc.nci.nih.gov/>.  
AUTHORS 1 (bases 1 to 495)  
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished (1999)  
CONTACT: Robert Strausberg, Ph.D.  
Email: [cgabbs-remail.nih.gov](mailto:cgabbs-remail.nih.gov)  
Tissue Procurement: The Cepko Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
Plate: L1AM1930 row: f column: 07  
High quality sequence stop: 489.  
Location/Qualifiers

FEATURES  
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/organism="Mus musculus"  
/db\_xref="taxon:10090"

/clone="IMAGE:5365638"  
/clone\_lib="NIH\_MGC\_94"  
/tissue\_type="retina"  
/lab\_host="DH10B (Phage-resistant)"  
/note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI; Site\_2: SalI; Cloned unidirectionally; oligo-dT primed. Average insert size 3.3 kb. Library enriched for full-length clones and constructed by Life Technologies. Note: this is a NIH\_MGC library."

BASE COUNT 105 a 159 c 111 g 120 t

Query Match 93.6%; Score 23.4; DB 13; Length 495;  
Best Local Similarity 96.0%; Pred. No. 21;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTGTGAGAGCTCTTCCA 25  
|||||  
Db 270 GCTTCTTGTGAGAGCTCTTCCA 294

RESULT 3 628 bp mRNA linear EST 20-SEP-2001  
LOCUS B1734160 603351436F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:5358776 5',  
DEFINITION mRNA sequence.  
ACCESSION B1734160  
VERSION B1734160.1 GI:15711173  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Mammalia; Chordata; Craniata; Vertebrata; Euteleostomi; Eukaryota; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE NIH-MGC <http://mgc.nci.nih.gov/>.  
AUTHORS 1 (bases 1 to 628)  
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished (1999)  
CONTACT: Robert Strausberg, Ph.D.  
Email: [cgabbs-remail.nih.gov](mailto:cgabbs-remail.nih.gov)  
Tissue Procurement: The Cepko Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
Plate: L1AM1912 row: h column: 09  
High quality sequence start: 4  
High quality sequence stop: 627.  
Location/Qualifiers

FEATURES  
Source 1. 628  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:5358776"  
/clone\_lib="NIH\_MGC\_94"  
/tissue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI; Site\_2: SalI; Cloned unidirectionally; oligo-dT primed. Average insert size 3.3 kb. Library enriched for full-length clones and constructed by Life Technologies. Note: this is a NIH\_MGC library."

BASE COUNT 159 a 180 c 152 g 137 t

Query Match 93.6%; Score 23.4; DB 13; Length 628;  
Best Local Similarity 96.0%; Pred. No. 23;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTGTGAGAGCTCTTCCA 25  
|||||  
Db 45 GCTTCTTGTGAGAGCTCTTCCA 69

RESULT 4  
 LOCUS B1736587  
 DEFINITION 603361053F1 NIH\_MGC\_94 Mus musculus CDNA clone IMAGE:536804 5',  
 mRNA sequence.  
 ACCESSION B1736587  
 VERSION B1736587.1 GI:15713600  
 KEYWORDS EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 REFERENCE NIH-MGC http://mgi.nci.nih.gov/.  
 1 (bases 1 to 636)  
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
 JOURNAL Unpublished (1999)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgaabs-remail.nih.gov  
 Tissue Procurement: The Cepko Laboratory  
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LLAM1936 row: 3 column: 13  
 High quality sequence stop: 632.  
 Location/Qualifiers  
 1..636  
 /organism="Mus musculus"  
 /db\_xref="taxon:10090"  
 /clone="IMAGE:536804"  
 /clone\_1ib="NIH\_MGC\_94"  
 /tissue\_type="retina"  
 /lab\_host="DH10B (phage-resistant)"  
 /note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI;  
 Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
 Average insert size 3.3 kb. Library enriched for  
 full-length clones and constructed by Life Technologies.  
 Note: this is a NIH\_MGC Library."  
 BASE COUNT 145 a 208 c 147 g 136 t  
 ORIGIN  
 Query Match 93.6%; Score 23.4; DB 13; Length 636;  
 Best Local Similarity 96.0%; Pred. No. 24;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 GCTTCTTTGCTGAGAGCTCTTCCA 25  
 |||||||||  
 Db 179 GCTTCTTTGCTGAGAGCTCTTCCA 203  
 RESULT 5  
 LOCUS BG403992  
 DEFINITION 602419859F1 NIH\_MGC\_94 Mus musculus CDNA clone IMAGE:4526966 5',  
 mRNA sequence.  
 ACCESSION BG403992  
 VERSION BG403992.1 GI:13297440  
 KEYWORDS EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 REFERENCE NIH-MGC http://mgi.nci.nih.gov/.  
 1 (bases 1 to 649)  
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
 JOURNAL Unpublished (1999)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgaabs-remail.nih.gov  
 Tissue Procurement: The Cepko Laboratory  
 CDNA Library Preparation: Life Technologies, Inc.

CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LLAM10435 row: e column: 15  
 High quality sequence stop: 644.  
 Location/Qualifiers  
 1..649  
 /organism="Mus musculus"  
 /db\_xref="taxon:10090"  
 /clone="IMAGE:4526966"  
 /clone\_1ib="NIH\_MGC\_94"  
 /tissue\_type="retina"  
 /lab\_host="DH10B (phage-resistant)"  
 /note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI;  
 Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
 Average insert size 3.3 kb. Library enriched for  
 full-length clones and constructed by Life Technologies.  
 Note: this is a NIH\_MGC Library."  
 BASE COUNT 164 a 203 c 140 g 142 t  
 ORIGIN  
 Query Match 93.6%; Score 23.4; DB 12; Length 649;  
 Best Local Similarity 96.0%; Pred. No. 24;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 GCTTCTTTGCTGAGAGCTCTTCCA 25  
 |||||||||  
 Db 108 GCTTCTTTGCTGAGAGCTCTTCCA 132  
 RESULT 6  
 LOCUS B1730233  
 DEFINITION B1730233 651 bp mRNA linear EST 20-SEP-2001  
 60335020F1 NIH\_MGC\_94 Mus musculus CDNA clone IMAGE:5357472 5',  
 mRNA sequence.  
 ACCESSION B1730233  
 VERSION B1730233.1 GI:15707246  
 KEYWORDS EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 REFERENCE NIH-MGC http://mgi.nci.nih.gov/.  
 1 (bases 1 to 651)  
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
 JOURNAL Unpublished (1999)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgaabs-remail.nih.gov  
 Tissue Procurement: The Cepko Laboratory  
 CDNA Library Preparation: Life Technologies, Inc.  
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LLAM1909 row: b column: 01  
 High quality sequence stop: 539.  
 Location/Qualifiers  
 1..651  
 /organism="Mus musculus"  
 /db\_xref="taxon:10090"  
 /clone="IMAGE:5357472"  
 /clone\_1ib="NIH\_MGC\_94"  
 /tissue\_type="retina"  
 /lab\_host="DH10B (phage-resistant)"  
 /note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI;  
 Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
 Average insert size 3.3 kb. Library enriched for  
 full-length clones and constructed by Life Technologies.  
 Note: this is a NIH\_MGC Library."  
 BASE COUNT 154 a 198 c 149 g 150 t

ORIGIN

Query Match 93.6%; Score 23.4; DB 13; Length 651;  
 Best Local Similarity 96.0%; Pred. No. 24;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTGGCTGAGAGCTCTTCCA 25  
 |||||  
 Db 61 GCTTCTTGGCTGAGAGCTCTTCCA 85

RESULT 7  
 B1730180 656 bp mRNA linear EST 20-SEP-2001  
 LOCUS 603349724F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:5357429 5',  
 DEFINITION mRNA sequence.  
 B1730180  
 VERSION B1730180.1 GI:15707193  
 KEYWORDS EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus

REFERENCE  
 AUTHORS Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 TITLE NIH-MGC http://mgi.nci.nih.gov/.  
 JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)  
 COMMENT Unpublished (1999)  
 Contact: Robert Strausberg, Ph.D.  
 Email: cgaps-remail.nih.gov  
 Tissue Procurement: The Cepko Laboratory  
 cDNA Library Preparation: Life Technologies, Inc.  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LLAM1908 row: p column: 06  
 High quality sequence stop: 649.

FEATURES  
 source Location/Qualifiers  
 1..656  
 /organism="Mus musculus"  
 /db\_xref="taxon:10090"  
 /clone\_image="5357429"  
 /clone\_lib="NIH\_MGC\_94"  
 /tissue\_type="retina"  
 /lab\_host="DH10B (phage-resistant)"  
 /note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI;  
 Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
 Average insert size 3.3 kb. Library enriched for  
 full-length clones and constructed by Life Technologies.  
 Note: this is a NIH\_MGC Library."

BASE COUNT 142 a 210 c 152 g 152 t

ORIGIN

Query Match 93.6%; Score 23.4; DB 13; Length 656;  
 Best Local Similarity 96.0%; Pred. No. 24;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTGGCTGAGAGCTCTTCCA 25  
 |||||  
 Db 402 GCTTCTTGGCTGAGAGCTCTTCCA 426

RESULT 8  
 B1729678 663 bp mRNA linear EST 20-SEP-2001  
 LOCUS 603349362F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:5356959 5',  
 DEFINITION mRNA sequence.  
 B1729678  
 VERSION B1729678.1 GI:15706691  
 KEYWORDS EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus

REFERENCE  
 AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 TITLE Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 JOURNAL NIH-MGC http://mgi.nci.nih.gov/.  
 COMMENT Unpublished (1999)  
 Contact: Robert Strausberg, Ph.D.  
 Email: cgaps-remail.nih.gov  
 Tissue Procurement: The Cepko Laboratory  
 cDNA Library Preparation: Life Technologies, Inc.  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LLAM1907 row: 1 column: 16  
 High quality sequence stop: 659.

FEATURES  
 source Location/Qualifiers  
 1..663  
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 /db\_xref="taxon:10090"  
 /clone\_image="5356959"  
 /clone\_lib="NIH\_MGC\_94"  
 /tissue\_type="retina"  
 /lab\_host="DH10B (phage-resistant)"  
 /note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI;  
 Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
 Average insert size 3.3 kb. Library enriched for  
 full-length clones and constructed by Life Technologies.  
 Note: this is a NIH\_MGC Library."

BASE COUNT 156 a 209 c 151 g 147 t

ORIGIN

Query Match 93.6%; Score 23.4; DB 13; Length 663;  
 Best Local Similarity 96.0%; Pred. No. 24;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTGGCTGAGAGCTCTTCCA 25  
 |||||  
 Db 152 GCTTCTTGGCTGAGAGCTCTTCCA 176

RESULT 9  
 BG298553 672 bp mRNA linear EST 21-FEB-2001  
 LOCUS BG298553  
 DEFINITION 602336937F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:4511697 5',  
 mRNA sequence.  
 BG298553  
 VERSION BG298553.1 GI:13063322  
 KEYWORDS EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus

REFERENCE  
 AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 TITLE Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 JOURNAL NIH-MGC http://mgi.nci.nih.gov/.  
 COMMENT Unpublished (1999)  
 Contact: Robert Strausberg, Ph.D.  
 Email: cgaps-remail.nih.gov  
 Tissue Procurement: The Cepko Laboratory  
 cDNA Library Preparation: Life Technologies, Inc.  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LLAM0395 row: 1 column: 10  
 High quality sequence stop: 666.

FEATURES  
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 /db\_xref="taxon:10090"



/clone="IMAGE:451697"  
/clone.lib="NIH\_MGC\_94"  
/issue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: PCMV-SPORT6; Site:1: NotI;  
Site:2: SalI; Cloned unidirectionally; oligo-dt primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."  
BASE COUNT 150 a 206 c 156 g 160 t

Query Match 93.6%; Score 23.4; DB 12; Length 672;  
Best Local Similarity 96.0%; Pred. No. 24;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTCTGAGAGCTCTTCCA 25  
|||||  
DB 341 GCTTCTTCTGAGAGCTCTTCCA 365

RESULT 10  
BI736996 676 bp mRNA linear EST 20-SEP-2001  
LOCUS 603360842F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:5367875 5',  
DEFINITION mRNA sequence.  
BI736996  
ACCESSION BI736996  
VERSION BI736996.1 GI:15714009  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 676)  
AUTHORS NIH-MGC http://mgc.nci.nih.gov/  
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished (1999)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgabbs-remail.nih.gov  
Tissue Procurement: The Cepko Laboratory

cDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNL at:  
http://image.llnl.gov  
Plate: LHAM1936 row: c column: 12  
High quality sequence stop: 662.  
Location/Qualifiers

## FEATURES

source  
1. 676  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:5367875"  
/clone.lib="NIH\_MGC\_94"  
/issue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: PCMV-SPORT6; Site:1: NotI;  
Site:2: SalI; Cloned unidirectionally; oligo-dt primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."

## BASE COUNT

ORIGIN 144 a 214 c 160 g 158 t

Query Match 93.6%; Score 23.4; DB 13; Length 676;  
Best Local Similarity 96.0%; Pred. No. 24;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTCTGAGAGCTCTTCCA 25  
|||||  
DB 409 GCTTCTTCTGAGAGCTCTTCCA 433

RESULT 11  
BI735044 681 bp mRNA linear EST 20-SEP-2001  
LOCUS 603356176F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:5363395 5',  
DEFINITION mRNA sequence.  
BI735044  
ACCESSION BI735044  
VERSION BI735044.1 GI:15712057  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 681)  
AUTHORS NIH-MGC http://mgc.nci.nih.gov/  
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished (1999)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgabbs-remail.nih.gov  
Tissue Procurement: The Cepko Laboratory

cDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNL at:  
http://image.llnl.gov  
Plate: LHAM1924 row: h column: 20  
High quality sequence stop: 680.  
Location/Qualifiers

## FEATURES

source  
1. 681  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:5363395"  
/clone.lib="NIH\_MGC\_94"  
/issue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: PCMV-SPORT6; Site:1: NotI;  
Site:2: SalI; Cloned unidirectionally; oligo-dt primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH\_MGC Library."

BASE COUNT 152 a 220 c 156 g 152 t 1 others

Query Match 93.6%; Score 23.4; DB 13; Length 681;  
Best Local Similarity 96.0%; Pred. No. 24;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTCTTCTGAGAGCTCTTCCA 25  
|||||  
DB 244 GCTTCTTCTGAGAGCTCTTCCA 268

## RESULT 12

BI735809 682 bp mRNA linear EST 21-FEB-2001  
LOCUS 602939229F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:4505134 5',  
DEFINITION mRNA sequence.  
BI735809  
ACCESSION BI735809  
VERSION BI735809.1 GI:13057815  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 682)  
AUTHORS NIH-MGC http://mgc.nci.nih.gov/  
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished (1999)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgabbs-remail.nih.gov  
Tissue Procurement: The Cepko Laboratory

cDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)

DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>  
 Place: LLAM10378 row: g column: 23  
 High quality sequence stop: 603.  
 Location/Qualifiers

## FEATURES

source

1..682  
 /organism="Mus musculus"  
 /db\_xref="taxon:10090"  
 /clone="IMAGE:4505134"  
 /clone\_1lb="NIH\_MGC\_94"  
 /tissue\_type="retina"  
 /lab\_host="DH10B (phage-resistant)"  
 /note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI;  
 Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
 Average insert size 3.3 kb. Library enriched for  
 full-length clones and constructed by Life Technologies.  
 Note: this is a NIH\_MGC library."

BASE COUNT 137 a 205 c 167 g 173 t

## ORIGIN

Query Match 93.6%; Score 23.4; DB 12; Length 682;  
 Best Local Similarity 96.0%; Pred. No. 24;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTTGCTGAGAGCTCTTCCA 25

Db 512 GCTTCTTTGCTGAGAGCTCTTCCA 536

## RESULT 13

LOCUS

B1737008 694 bp mRNA linear EST 20-SEP-2001  
 DEFINITION 60336085F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:5368117 5',  
 mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

house mouse.  
 Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus;  
 1 (bases 1 to 694)  
 NIH-MGC <http://mgc.nci.nih.gov/>

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Unpublished (1999)  
 National Institutes of Health, Mammalian Gene Collection (MGC)  
 Contact: Robert Strausberg, Ph.D.  
 Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)

Tissue Procurement: The Cepko Laboratory  
 cDNA Library Preparation: Life Technologies, Inc.

DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>

Plate: LLAM11936 row: m column: 14  
 High quality sequence stop: 694.  
 Location/Qualifiers

## FEATURES

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 /db\_xref="taxon:10090"  
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 /tissue\_type="retina"  
 /lab\_host="DH10B (phage-resistant)"  
 /note="Organ: eye; Vector: pCMV-SPORT6; Site\_1: NotI;  
 Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
 Average insert size 3.3 kb. Library enriched for  
 full-length clones and constructed by Life Technologies.  
 Note: this is a NIH\_MGC library."

## BASE COUNT

157 a 216 c 156 g 165 t

Query Match 93.6%; Score 23.4; DB 13; Length 694;  
 Best Local Similarity 96.0%; Pred. No. 24;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTTGCTGAGAGCTCTTCCA 25

Db 363 GCTTCTTTGCTGAGAGCTCTTCCA 387

## RESULT 14

LOCUS

BG298167 696 bp mRNA linear EST 21-FEB-2001  
 DEFINITION 60239630F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:4507824 5',  
 mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

house mouse.  
 Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus;  
 1 (bases 1 to 696)  
 NIH-MGC <http://mgc.nci.nih.gov/>

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Unpublished (1999)  
 National Institutes of Health, Mammalian Gene Collection (MGC)  
 Contact: Robert Strausberg, Ph.D.  
 Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)

Tissue Procurement: The Cepko Laboratory  
 cDNA Library Preparation: Life Technologies, Inc.  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>

Plate: LLAM10385 row: h column: 01  
 High quality sequence stop: 677.  
 Location/Qualifiers

## FEATURES

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 Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
 Average insert size 3.3 kb. Library enriched for  
 full-length clones and constructed by Life Technologies.  
 Note: this is a NIH\_MGC library."

## BASE COUNT

148 a 218 c 164 g 166 t

## ORIGIN

Query Match 93.6%; Score 23.4; DB 12; Length 696;  
 Best Local Similarity 96.0%; Pred. No. 24;  
 Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 GCTTCTTTGCTGAGAGCTCTTCCA 25

Db 459 GCTTCTTTGCTGAGAGCTCTTCCA 483

## RESULT 15

LOCUS

B1735675 697 bp mRNA linear EST 20-SEP-2001  
 DEFINITION 60335782F1 NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:5364964 5',  
 mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

house mouse.  
 Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

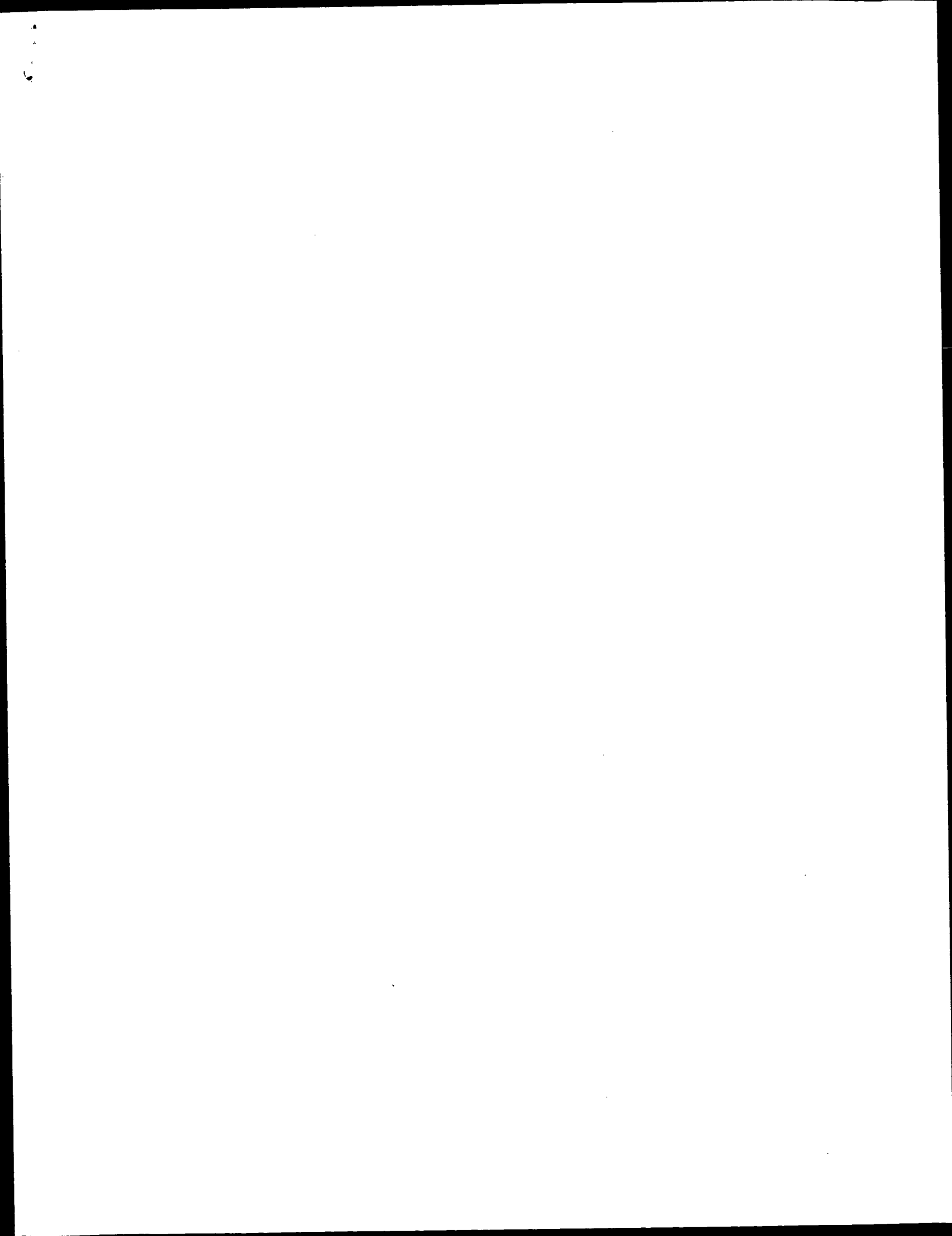
Mammalia; Euthera; Rodentia; Sciurognath1; Muridae; Murinae; Mus.  
1 (bases 1 to 697)  
NIH-MGC <http://mgc.ncl.nih.gov/>.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
Tissue Procurement: The Cepko Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>  
Plate: LLAM11928 row: 1 column: 05  
High quality sequence stop: 697.

FEATURES  
SOURCE

Location/Qualifiers  
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/tissue\_type="retina"  
/lab\_host="DH10B (phage-resistant)"  
/note="Organ: eye; Vector: PCMV-SPORT6; Site\_1: NotI;  
Site\_2: SalI; Cloned unidirectionally; oligo-dT primed.  
Average insert size 3.3 kb. Library enriched for  
full-length clones and constructed by Life Technologies.  
Note: this is a NIH-MGC Library."  
BASE COUNT 136 a 209 c 168 g 184 t  
ORIGIN

Query Match 93.6%; Score 23.4; DB 13; Length 697;  
Best Local Similarity 96.0%; Pred. No. 24;  
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
OY 1 GCTTCTTGCTGAGACTCTCA 25  
|||||||  
DB 585 GCTTCTTGCTGAGACTCTCA 609

Search completed: March 17, 2003, 13:09:23  
Job time : 886.737 secs



GenCore version 5.1.4.p5.4578  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 09:43:42 ; Search time 202.495 Seconds

(without alignments)  
3161.870 Million cell updates/sec

Title: US-09-836-439-6

Perfect score: 22

Sequence: 1 aagactctgagtaacaa 22

Scoring table: IDENTITY\_NUC

Gapop 10.0, Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : GenEmbl.\*

1: gb\_da.\*

2: gb\_hlg.\*

3: gb\_in.\*

4: gb\_cm.\*

5: gb\_ov.\*

6: gb\_pat.\*

7: gb\_ph.\*

8: gb\_pl.\*

9: gb\_pr.\*

10: gb\_ro.\*

11: gb\_sts.\*

12: gb\_sy.\*

13: gb\_un.\*

14: gb\_vl.\*

15: em\_da.\*

16: em\_fun.\*

17: em\_hum.\*

18: em\_in.\*

19: em\_mu.\*

20: em\_om.\*

21: em\_or.\*

22: em\_ov.\*

23: em\_pat.\*

24: em\_ph.\*

25: em\_pl.\*

26: em\_ro.\*

27: em\_sts.\*

28: em\_un.\*

29: em\_vl.\*

30: em\_hlg\_hum.\*

31: em\_hlg\_inv.\*

32: em\_hlg\_other.\*

33: em\_hlg\_mus.\*

34: em\_hlg\_pln.\*

35: em\_hlg\_rod.\*

36: em\_hlg\_mam.\*

37: em\_hlg\_vrt.\*

38: em\_sv.\*

39: em\_hlgo\_hum.\*

40: em\_hlgo\_mus.\*

41: em\_hlgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19.4	88.2	17938	AF198437	AF198437 Thelateria
2	18.8	85.5	42208	AL589989	AL589989 Human DNA
3	18.8	85.5	116685	HS516C23	293021 Human DNA s
4	18.8	85.5	178168	AC012350	AC012350 Homo sapi
5	18.8	85.5	180970	AC009302	AC009302 Homo sapi
6	18.8	85.5	182152	AC067929	AC067929 Homo sapi
7	18.8	85.5	189281	AC012458	AC012458 Homo sapi
8	18.8	85.5	191434	AC087477	AC087477 Homo sapi
9	18.8	85.5	197437	AL772303	AL772303 Homo sapi
10	18.8	85.5	223761	AC127239	AC127239 Mus muscu
11	18.8	85.5	229247	AC098888	AC098888 Mus muscu
12	18.8	85.5	317515	AC099415	AC099415 Mus muscu
13	18.4	83.6	10243	AE011043	AE011043 Methanosa
14	18.4	83.6	10467	AE010860	AE010860 Methanosa
15	18.4	83.6	152556	CNS01DSY	AL122035 Human chr
16	18.4	83.6	153951	AC019027	AC019027 Homo sapi
17	18.4	81.8	148507	AC011263	AC011263 Homo sapi
18	18.4	81.8	158633	AC015472	AC015472 Homo sapi
19	18.4	81.8	178199	AC018772	AC018772 Homo sapi
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21	17.8	80.9	907	AF213436	AF213436 Homo sapi
22	17.8	80.9	927	107695	107695 Sequence 18
23	17.8	80.9	3159	HS05SE1	X51934 Human dyslr
24	17.8	80.9	57307	AL603652	AL603652 Human DNA
25	17.8	80.9	83007	AC097448	AC097448 Homo sapi
26	17.8	80.9	107406	AC013491	AC013491 Homo sapi
27	17.8	80.9	141984	F91L	AC007591 Arabidops
28	17.8	80.9	148571	AC105316	AC105316 Homo sapi
29	17.8	80.9	161874	AC079864	AC079864 Homo sapi
30	17.8	80.9	163132	AC097180	AC097180 Ratcus no
31	17.8	80.9	176287	AL513347	AL513347 Mouse DNA
32	17.8	80.9	179591	AC093902	AC093902 Homo sapi
33	17.8	80.9	182463	AC120773	AC120773 Ratcus no
34	17.8	80.9	182933	AC097610	AC097610 Ratcus no
35	17.8	80.9	184573	AC111144	AC111144 Mus muscu
36	17.8	80.9	191664	AC125751	AC125751 Ratcus no
37	17.8	80.9	200539	AC121805	AC121805 Mus muscu
38	17.8	80.9	201296	AC115299	AC115299 Mus muscu
39	17.8	80.9	203753	AC104343	AC104343 Homo sapi
40	17.8	80.9	207945	AC117841	AC117841 Ratcus no
41	17.8	80.9	207974	AL831771	AL831771 Mus muscu
42	17.8	80.9	208953	CNS01MR3	AL160314 Human chr
43	17.8	80.9	210779	AC007450	AC007450 Homo sapi
44	17.4	79.1	12024	AE000995	AE000995 Archaeogl
45	17.4	79.1	65499	AC100433	AC100433 Mus muscu

## ALIGNMENTS

RESULT 1

AF198437

LOCUS

DEFINITION

AF198437 17938 bp DNA linear INV 05-DEC-2000

Thelateria parva strain Muguga hypothetical telomeric SfiI fragment

20 protein 3, hypothetical telomeric SfiI fragment 20 protein 2,

and hypothetical telomeric SfiI fragment 20 protein 1 genes,

complete cds.

AF198437 L36964

AF198437.1 GI:11545211

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Thelateria parva.

Eukaryota; Alveolata; Apicomplexa; Piroplasmida; Thelateridae;

REFERENCE

1 (bases 17114 to 17938)

AUTHORS Sohanpal,B.K., Morzaria,S.P., Gobright,E.I. and Bishop,R.P.  
TITLE Characterisation of the telomeres at opposite ends of a 3 Mb  
Theileria parva chromosome  
JOURNAL Nucleic Acids Res. 23 (11), 1942-1947 (1995)  
MEDLINE 95319937  
PUBMED 7596822  
REFERENCE 2 (bases 1 to 17938)  
AUTHORS Bishop,R., Gobright,E., Nene,V., Morzaria,S., Musoke,A. and  
Sohanpal,B.  
TITLE Polymorphic open reading frames encoding secretory proteins are  
located less than 3 kilobases from Theileria parva telomeres  
JOURNAL Mol. Biochem. Parasitol. 110 (2), 359-371 (2000)  
MEDLINE 20520968  
PUBMED 11071289  
REFERENCE 3 (bases 17114 to 17938)  
AUTHORS Sohanpal,B.K., Morzaria,S.P., Gobright,E.I. and Bishop,R.P.  
TITLE Direct Submission  
JOURNAL Submitted (16-JAN-1995) Unit 5, International Livestock Research  
Institute (ILRI), Old Naivasha Road, Nairobi P.O. Box 30709, Kenya  
4 (bases 1 to 17938)  
REFERENCE Bishop,R.P., Gobright,E.I. and Sohanpal,B.K.  
TITLE Direct Submission  
JOURNAL Submitted (25-OCT-1999) Unit 5, International Livestock Research  
Institute (ILRI), Old Naivasha Road, Nairobi P.O. Box 30709, Kenya  
COMMENT On Dec 5, 2000 this sequence version replaced gi:624245.  
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CDS

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CDS

CDS

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CDS

repeat\_region

BASE COUNT 5477 a 3549 c 3098 g 5814 t  
ORIGIN  
Query Match 88.2%; Score 19.4; DB 3; Length 17938;

Best Local Similarity 95.2%; Pred. No. 56;  
 Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY      1 AAGACTTCTGAGTAACAATCA 21
      || ||||| ||||| ||||| |||||
Db      8185 AATACCTCTGAGTAACAATCA 8205
  
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RESULT 2	
AL589989	
LOCUS	
DEFINITION	AL589989 42208 bp DNA linear PRI 12-MAY-2001
ACCESSION	Human DNA sequence from clone RP1-116B8 on chromosome 6, complete sequence. ATG00000

ACCESSION	AL268385	GI:14041769
VERSION	AL589989.6	
KEYWORDS	HTG.	
SOURCE	human.	
ORGANISM	Homo sapiens	

**REFERENCE**                    **"(bases 1 to 42208)**  
**AUTHORS**                    **Tromans, A.**  
**TITLE**                         **Direct Submission**  
**JOURNAL**                      **Submitted (12-MAY-2001) Sanger Centre, Hinxton, Cambridgeshire,**

**COMMENT**

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction fragment digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em', EMBL; Sw', SWISSPROT; Tr', TREMBL; Wp', WORMPEP; Information on the WORMPEP database can be found at

<http://www.sanger.ac.uk/Projects/Celegans/wormpep> This sequence was generated from part of bacterial clone contl15 of human chromosome 6, constructed by the Sanger Centre Human Mapping Group. Further information can be found at

<http://www.sanger.ac.uk/HGP/Chr6>

RP1-11688 is from the library RGC1-1 constructed by the group of Pieter de Jong. For further details see

<http://www.chori.org/Dacpac/home.htm>

VECTOR: pCYPAC2

IMPORTANT: This sequence is not the entire insert of clone RP1-11688 it may be shorter because we sequence overlapping sections only once, except for the 100 base overlap.

The true left end of clone RP1-13115 is at 42105 in this sequence. The true right end of clone RP3-48313 is at 100 in this sequence.

insert:42000-11688

**FEATURES**  
**SOURCE**

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2339. .2368
repeat_region
/note="15 copies 2 mer tt 86% conserved"
3125. .3190
repeat_region
/note="33 copies 2 mer ca 66% conserved"

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repeat_region	3464. .3770	/note="AluV repeat: matches 1. .309 of consensus"
repeat_region	3996. .4294	/note="AluSx repeat: matches 1. .300 of consensus"
repeat_region	4295. .4364	/note="MER4A repeat: matches 655. .728 of consensus"
repeat_region	4374. .6566	/note="TIGER1 repeat: matches 2. .2227 of consensus"
repeat_region	6823. .7007	/note="AluV repeat: matches 116. .307 of consensus"
repeat_region	7013. .7190	/note="TIGER1 repeat: matches 2241. .2418 of consensus"
repeat_region	7450. .7711	/note="MER4A repeat: matches 2. .238 of consensus"
repeat_region	7988. .8091	/note="MIR repeat: matches 41. .146 of consensus"
repeat_region	8780. .8964	/note="L1M49 repeat: matches 6087. .6270 of consensus"
repeat_region	9608. .9657	/note="AluJ/FLM repeat: matches 10. .59 of consensus"
repeat_region	10806. .10945	/note="MIR repeat: matches 13. .148 of consensus"
repeat_region	11078. .11372	/note="AluJo repeat: matches 4. .298 of consensus"
repeat_region	12599. .13021	/note="L1MB3 repeat: matches 5737. .6184 of consensus"
repeat_region	13039. .13314	/note="AluJo repeat: matches 7. .298 of consensus"
repeat_region	14353. .14411	/note="L2 repeat: matches 2405. .2459 of consensus"
repeat_region	14412. .14724	/note="AluJo repeat: matches 1. .301 of consensus"
repeat_region	14725. .14934	/note="L2 repeat: matches 2459. .2691 of consensus"
repeat_region	16429. .16820	/note="MER63A repeat: matches 44. .208 of consensus"
repeat_region	17156. .17192	/note="MER63A repeat: matches 1. .37 of consensus"
repeat_region	17998. .18335	/note="L2 repeat: matches 2158. .2506 of consensus"
repeat_region	18384. .18689	/note="MER2 repeat: matches 22. .345 of consensus"
repeat_region	18690. .19270	/note="L1MB3 repeat: matches 5434. .6176 of consensus"
repeat_region	22150. .22336	/note="L2 repeat: matches 1246. .1462 of consensus"
repeat_region	22396. .22600	/note="L2 repeat: matches 112. .316 of consensus"
repeat_region	22706. .22754	/note="MER3 repeat: matches 152. .201 of consensus"
repeat_region	22755. .22883	/note="FLM4A repeat: matches 1. .130 of consensus"
repeat_region	22884. .23038	/note="MER3 repeat: matches 3. .152 of consensus"
repeat_region	23345. .23433	/note="L1P43 repeat: matches 6063. .6156 of consensus"
repeat_region	23561. .23632	/note="36 copies 2 mer at 65% conserved"
repeat_region	23694. .23989	/note="MIR1A1 repeat: matches 44. .315 of consensus"
repeat_region	24062. .24213	/note="MER5B repeat: matches 3. .173 of consensus"
repeat_region	25110. .25213	/note="MIR repeat: matches 51. .157 of consensus"
repeat_region	26282. .26447	/note="L2 repeat: matches 2482. .2626 of consensus"
repeat_region	26454. .26782	





repeat_region	/note="L1M1 repeat: matches 5398. .6314 of consensus 23343. .23754
repeat_region	/note="L2 repeat: matches 2029. .2520 of consensus 24063. .24264
repeat_region	/note="L1B repeat: matches 189. .390 of consensus 24302. .24345
repeat_region	/note="22 copies 2 mer tt 75% conserved" 24361. .30507
repeat_region	/note="L1P2 repeat: matches 4. .6144 of consensus 30510. .30679
repeat_region	/note="L1B1 repeat: matches 1. .170 of consensus 31272. .31782
repeat_region	/note="L2 repeat: matches 960. .2750 of consensus 31811. .32018
repeat_region	/note="M1R repeat: matches 39. .262 of consensus 32300. .33151
repeat_region	/note="L1B1 repeat: matches 5333. .6155 of consensus 33152. .33236
repeat_region	/note="L1S2P/9 repeat: matches 192. .276 of consensus 33484. .33868
repeat_region	/note="L2 repeat: matches 1862. .2267 of consensus 33887. .34010
repeat_region	/note="L1P3 repeat: matches 1. .143 of consensus 34187. .34218
repeat_region	/note="16 copies 2 mer aa 84% conserved" 34220. .34364
repeat_region	/note="L1P33 repeat: matches 382. .517 of consensus 34403. .34964
repeat_region	/note="L1M6 repeat: matches 272. .906 of consensus 35050. .35158
repeat_region	/note="L1M4 repeat: matches 6176. .6290 of consensus 35141. .35266
repeat_region	/note="L1M4 repeat: matches 2121. .2241 of consensus 35267. .35587
repeat_region	/note="A1U repeat: matches 1. .309 of consensus 35588. .36308
repeat_region	/note="L1M4 repeat: matches 2241. .2996 of consensus 36248. .36925
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repeat_region	/note="A1USq/x repeat: matches 1. .129 of consensus 37878. .37923
repeat_region	/note="23 copies 2 mer ca 100% conserved" 37928. .38256
repeat_region	/note="A1USx repeat: matches 1. .300 of consensus 38262. .38433
repeat_region	/note="L1M4 repeat: matches 3047. .3335 of consensus 38463. .38837
repeat_region	/note="L1M4 repeat: matches 3790. .4186 of consensus 38847. .39138
repeat_region	/note="A1UJO repeat: matches 12. .293 of consensus 39344. .39389
repeat_region	/note="23 copies 2 mer aa 76% conserved" 40977. .41071
repeat_region	/note="M1R repeat: matches 12. .120 of consensus 41529. .41819
repeat_region	/note="L2 repeat: matches 1780. .2071 of consensus 41800. .41805
misc_feature	/note="152 transposable element excised from this position" 41820. .42230
repeat_region	/note="M1T1J repeat: matches 56. .505 of consensus 42267. .42627
repeat_region	/note="L2 repeat: matches 2301. .2677 of consensus 42644. .42731
repeat_region	/note="44 copies 2 mer aa 69% conserved" 43499. .43640
repeat_region	/note="71 copies 2 mer tt 63% conserved" 44092. .44158

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repeat_region /note="LTR1D repeat: matches 1. .66 of consensus"
44166. .44649
/note="LTR1D repeat: matches 5. .505 of consensus"
repeat_region 44686. .45053
/note="LTR1D repeat: matches 113. .503 of consensus"
repeat_region 45793. .45960
/note="MIR repeat: matches 7. .171 of consensus"
repeat_region 46079. .46271
/note="LTR1F repeat: matches 347. .541 of consensus"
repeat_region 46316. .46363
/note="LTR2CB repeat: matches 236. .283 of consensus"
repeat_region 46602. .46641
/note="20 copies 2 mer tg 95% conserved"
misc_feature complement(47028. .47443)
repeat_region /note="match: GSS B53031 clone 2008G6"
47509. .48290
/note="L1MB5 repeat: matches 535. .616 of consensus"
repeat_region 48283. .49518
/note="L1MB5 repeat: matches 3964. .5271 of consensus"
repeat_region 49519. .49586
/note="34 copies 2 mer ta 72% conserved"
repeat_region 49980. .50288
/note="AluSq repeat: matches 3. .311 of consensus"
repeat_region 50661. .51118
/note="L1MC4 repeat: matches 6102. .7977 of consensus"
misc_feature complement(51869. .52243)
repeat_region /note="match: GSS AQ109033 clone 2377A15"
52215. .52242
/note="14 copies 2 mer ac 100% conserved"
misc_feature <52243. .>52703
repeat_region /note="match: GSS AQ234004"
54296. .54852
/note="12 repeat: matches 2180. .2743 of consensus"
repeat_region 55942. .55097
/note="MIR repeat: matches 49. .210 of consensus"

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Query Match	85.5%	Score 18.8;	DB 9;	Length 116685;
Best Local Similarity	90.9%	Pred. NO. 90;		
Matches 20;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;

QY 1 AAGACTTCTGAGTAACAATCAA 22  
|| |||||  
Db 59346 AACTCTTCTGAGTAACAATCAA 59325

LOCUS 178168 bp DNA linear HTG 01-APR-2000  
 DEFINITION Homo sapiens clone Rpl1-16N9, WORKING DRAFT SEQUENCE, 16 unordered  
 pieces.  
 ACCESSION AC012350  
 VERSION AC012350.3 GI:7381803  
 KEYWORDS HTG; HTG\_\_PHASE1; HTG\_\_DRAFT.  
 SOURCE Homo sapiens.  
 ORGANISM Homo sapiens

REFERENCE	AUTHORS	TITLE	JOURNAL	REFERENCE
1 (bases 1 to 178168)	Blirren,B., Linton,L., Nusbaum,C. and Lander,E.	Homo sapiens, clone RP11-16N9	Unpublished	2 (bases 1 to 178168)
Blirren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,				

Bitten, B., Alnton, L., Neudamm, C., Lander, E., Allen, N., Anderson, M.,  
 Baldwin, J., Barua, N., Beckerly, R., Boguslavsky, L., Bourkpatrick, B.,  
 Brown, A., Castle, A., Colangelo, M., Collins, S., Collymore, A.,  
 Cooke, P., Deardellano, K., Dewar, K., Domingo, M., Donelan, L., Doyle, M.,  
 Ferrari, P., Fitzhugh, W., Forrest, C., Funke, R., Gage, D., Gallegan, J.,  
 Galarza, J., Gadyana, S., Grant, G., Hagos, B., Heatford, A., Horton, L.,  
 Howland, J. C., Johnson, R., Jones, C., Kann, L., Karatas, A., Klein, J.,  
 Lehoczy, J., Lieu, C., Locke, K., Macdonald, P., Marquis, N.,  
 McEwan, P., McGurk, A., McKernan, K., McLaughlin, J., Meldrum, J.,  
 Morrow, K., Naylor, J., Norman, C. H., O'Connor, T., O'Donnell, P.,  
 Peterson, K., Pollara, V., Riley, R., Roy, A., Santos, R., Severy, P.,

TITLE  
JOURNAL  
COMMENT

Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J.,  
Tesfaye, S., Tirrell, A., Vassiliev, H., Vo, A., Wheeler, J., Wu, X.,  
Wyman, D., Ye, W.-J., Zimmer, A. and Zody, M.  
Direct Submission  
Submitted (25-OCT-1999) Whitehead Institute/MIT Center for Genome  
Research, 320 Charles Street, Cambridge, MA 02141, USA  
On Apr 1, 2000 this sequence version replaced 91:6479001.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1997)  
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

## Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: MIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)

Project Information

Center project name: L3593

Center clone name: 16\_N-9

## Summary Statistics

Sequencing vector: M13; M77815; 100% of reads  
Chemistry: Dye-terminator Big Dye; 100% of reads  
Assembly program: Phrap; version 0.960731  
Consensus quality: 172070 bases at least Q40  
Consensus quality: 174479 bases at least Q30  
Consensus quality: 175597 bases at least Q20  
Insert size: 170000; agarose-gel  
Insert size: 176668; sum-of-contents  
Quality coverage: 6.4 in Q20 bases; agarose-gel  
Quality coverage: 6.2 in Q20 bases; sum-of-contents

\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 16 contigs. The true order in this sequence record is  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

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1 1694: contig of 1694 bp in length
1695 1794: gap of 100 bp
1795 4437: contig of 2643 bp in length
4438 4537: gap of 100 bp
4538 6644: contig of 2107 bp in length
6645 6744: gap of 100 bp
6745 10092: contig of 3348 bp in length
10093 10192: gap of 100 bp
10193 11801: contig of 1609 bp in length
11802 11901: gap of 100 bp
11902 15584: contig of 3683 bp in length
15585 15684: gap of 100 bp
15685 19675: contig of 3991 bp in length
19676 19775: gap of 100 bp
19776 27913: contig of 8138 bp in length
27914 28013: gap of 100 bp
28014 34156: contig of 6143 bp in length
34157 34256: gap of 100 bp
34257 43325: contig of 9069 bp in length
43326 43425: gap of 100 bp
43426 55265: contig of 11840 bp in length
55266 55365: gap of 100 bp
55366 68306: contig of 12941 bp in length
68307 68406: gap of 100 bp
68407 82910: contig of 14504 bp in length
82911 83010: gap of 100 bp
83011 102181: contig of 19171 bp in length
102182 102281: gap of 100 bp
102282 128056: contig of 25777 bp in length
128059 128158: gap of 100 bp
128159 178168: contig of 50010 bp in length.
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## FEATURES

## SOURCE

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/db\_xref="taxon:9606"

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1. 1694
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1795. 4437
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4538. 6644
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6745. 10092
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10193. 11801
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11902. 15584
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15685. 19675
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19776. 27913
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28014. 34156
/feature="assembly-fragment"
34257. 43325
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43426. 55265
/feature="assembly-fragment"
55366. 68306
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68407. 82910
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83011. 102181
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102282. 128058
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128159. 178168
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BASE COUNT 50059 a 39406 c 38562 g 48641 t 1500 others
ORIGIN
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Query Match 85.5%; Score 18.8; DB 2; Length 178168;
Best Local Similarity 90.9%; Pred. No. 86;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

1 ANGACTTGTGAGTACATCAA 22
16379 AACACTTGTGAGTACATCAA 16358
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RESULT 5
LOCUS AC009302 180970 bp DNA linear PRI 30-SEP-2000
DEFINITION Homo sapiens BAC clone RP11-71J24 from 2, complete sequence.
ACCESSION AC009302
VERSION AC009302.2 GI:9931944
KEYWORDS HTG.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 180970)
AUTHORS Sultson, J.E. and Waterston, R.
Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE 99063792
PUBMED 9847074
REFERENCE 2 (bases 1 to 180970)
AUTHORS Tomlinson, C., Wohldmann, P., Maupin, R. and Reitz, L.
The sequence of Homo sapiens BAC clone RP11-71J24
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 180970)
AUTHORS Waterston, R.H.
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TITLE Direct Submission  
JOURNAL Submitted (13-AUG-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA  
REFERENCE 4 (bases 1 to 180970)  
AUTHORS Waterston, R.H.  
TITLE Direct Submission  
JOURNAL Submitted (28-AUG-2000) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA  
REFERENCE 5 (bases 1 to 180970)  
AUTHORS Waterston, R.H.  
TITLE Direct Submission  
JOURNAL Submitted (10-SEP-2000) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA  
REFERENCE 6 (bases 1 to 180970)  
AUTHORS Waterston, R.  
TITLE Direct Submission  
JOURNAL Submitted (30-SEP-2000) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA  
COMMENT On Aug 28, 2000 this sequence version replaced gl:5732171.  
----- Genome Center  
Center: Washington University Genome Sequencing Center  
Center code: WUGSC  
Web site: <http://genome.wustl.edu/gsc>  
Contact: saplens@wustl.wustl.edu  
----- Summary Statistics  
Center project name: H\_NH0071J24  
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NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

#### MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

#### SOURCE INFORMATION:

The RPCT-11 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Woon, P. Y., Zhao, B., Frenken, E., Tateo, M., Catanesi, J. J., and de Jong, P. J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)  
VECTOR: pBAC3.6

#### NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-350124; the clone sequenced to the right is RP11-296A19. Actual start of this clone is at base position 1 of RP11-71J24; actual end is at base position 180970 of RP11-71J24.

#### FEATURES

##### source

Location/Qualifiers  
1..180970  
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/db\_xref="taxon:9606"  
/chromosome="2"  
/map="2"  
/clone="RP11-71J24"  
/clone\_1lb="RPCT-11"

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repeat_region	9014..9113	/rpt_family="L1"
repeat_region	9120..9530	/rpt_family="L1"
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Best local Similarity 90.9%; Pred. No. 85;
Matches 20; Conservativity 0; Mismatches 2; Indels 0; Gaps 0;

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Db 30424 AAGACTTCGAGCAAAATCAA 30445

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RESULT 6
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LOCUS Homo sapiens chromosome 6 clone RP11-403120 map 6, WORKING DRAFT
DEFINITION
AC067929
SEQUENCE, 18 unordered pieces.
AC067929
VERSION AC067929.2 GI:8247824
KEYWORDS HTG: HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 182152)
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 6, clone RP11-403120
unpublished
2 (bases 1 to 182152)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,
Boguslavsky,L., Boukhalter,B., Brown,A., Burkett,G.,
Campopiano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,
Collymore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
Dodge,S., Domino,M., Doyle,M., Ferreira,P., FitzHugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Grant,G., Hages,B., Heaford,A., Horton,L.,
Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,
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Levine,R., Liu,G., Locke,K., Macdonald,P., Markuis,N.,
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Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,
Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Testaye,S., Theodore,J., Tirrell,A., Travers,M., Triggillo,J.,
Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zimmer,A. and Zody,M.

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TITLE
JOURNAL
REFERENCE
AUTHORS

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TITLE
JOURNAL
COMMENT

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Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,
Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Testaye,S., Theodore,J., Tirrell,A., Travers,M., Triggillo,J.,
Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zimmer,A. and Zody,M.
Direct Submission
Submitted (27-APR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 182152)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,
Boguslavsky,L., Boukhalter,B., Brown,A., Burkett,G.,
Campopiano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,
Collymore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
Dodge,S., Domino,M., Doyle,M., Ferreira,P., FitzHugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Grant,G., Hages,B., Heaford,A., Horton,L.,
Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,
Klein,J., Lacroque,K., Lamazares,R., Landers,T., Lehoczy,J.,
Levine,R., Liu,G., Locke,K., Macdonald,P., Markuis,N.,
McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,
Meldrim,J., Meneus,L., Mihova,T., Miranda,C., Mienga,V., Morrow,J.,
Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,
Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Testaye,S., Theodore,J., Tirrell,A., Travers,M., Triggillo,J.,
Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zimmer,A. and Zody,M.
Direct Submission
Submitted (24-AUG-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jun 4, 2000 this sequence version replaced g1:7655991.
All repeats were identified using RepeatMasker:
Smit,A.F.A. & Green,P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: MIBR
Web site: http://www.seq.wi.mit.edu
Contract: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: 403_1-20
Center clone name: 403_1-20
----- Summary Statistics
Sequencing vector: M13; M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 173395 bases at least Q40
Consensus quality: 177417 bases at least Q30
Consensus quality: 179105 bases at least Q20
Insert size: 179000; agarose-fp
Insert size: 180452; sum-of-contigs
Quality coverage: 4.8 in Q20 bases; agarose-fp
Quality coverage: 4.7 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 2222: contig of 2222 bp in length
* 2223 2322: gap of 100 bp
* 2323 3785: contig of 1463 bp in length
* 3786 3885: gap of 100 bp
* 3886 6753: contig of 2866 bp in length

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*      9862      18746: contig of 8885 bp in length
*      18747      18846: gap of unknown length
*      18847      27396: contig of 8550 bp in length
*      27397      27496: gap of unknown length
*      27497      35489: contig of 7993 bp in length
*      35490      35589: gap of unknown length
*      35590      51472: contig of 15883 bp in length
*      51473      51572: gap of unknown length
*      51573      65229: contig of 13657 bp in length
*      65230      65329: gap of unknown length
*      65330      89666: contig of 24337 bp in length
*      89667      89766: gap of unknown length
*      89767      117943: contig of 28177 bp in length
*      117944      118044: gap of unknown length
*      118044      189281: contig of 71238 bp in length.
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*      4704. 9761
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*      misc_feature
*      27497. 35489
*      /note="assembly_name:Contig10"
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*      35590. 51472
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*      51573. 65229
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*      65330. 89666
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*      /note="assembly_name:Contig14"
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*      118044. 189281
*      /note="assembly_name:Contig15"
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*      54120 a 39374 c 39106 g 55769 t 912 others
*      BASE COUNT
*      ORIGIN
*      Query Match      85.5%; Score 18.8; DB 2; Length 189281;
*      Best Local Similarity 90.9%; Pred. No. 85;
*      Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
*      Oy      1 AAGACTTCTGAGTAAACATCAA 22
*      Db 181216 AAGACTTCTGAGTAAACATCAA 181237
*      RESULT 8
*      AC087477
*      LOCUS      AC087477      191434 bp      DNA      linear      PRI 16-JUL-2002
*      DEFINITION      Homo sapiens chromosome 15, clone RP11-522B15, complete sequence.
*      ACCESSION      AC087477
*      VERSION      AC087477.8      GI:21844630
*      KEYWORDS
*      HMG.
*      SOURCE
*      human.
*      Homo sapiens
*      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
*      Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
*      Birten,B., Nusbaum,C. and Lander,E.
*      1 (bases 1 to 191434)
*      TITLE
*      Homo sapiens chromosome 15, clone RP11-522B15
*      JOURNAL
*      Unpublished
*      REFERENCE
*      2 (bases 1 to 191434)

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AUTHORS
Birten,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B., Brown,A.,
Cammarata,J., Campoliano,A., Choepel,Y., Colangelo,M., Collins,S.,
Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
Dodge,S., Fato,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J.,
Gardyna,S., Ginde,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Hearford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Karatas,A., Larroque,K., Lamazares,R., Landers,T.,
Lehoczy,J., Levine,R., Liu,G., Maclean,C., Macdonald,P.,
Macpherson,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K.,
Meneus,L., Mlhova,T., Mlhova,T., Mlhova,T., Mlhova,V.,
Mphuy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Pollara,V., Pollara,V., Roman,J., Rosetti,M.,
Rogov,P., Roy,A., Santos,R., Roman,J., Rosetti,M.,
Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rosetti,M.,
Roy,A., Santos,R., Schauer,S., Schnupack,R., Seaman,S., Severy,P.,
Soungaz,C., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Straus,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J.,
Travers,M., Travis,N., Triggillo,J., Vassiliev,H., Viel,R., Vo,A.,
Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
TITLE
Direct Submission
JOURNAL
Submitted (03-JAN-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE
3 (bases 1 to 191434)
AUTHORS
Birten,B., Linton,L., Nusbaum,C., Lander,E., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L.,
Boukhgalter,B., Brown,A., Cammarata,J., Campoliano,A., Chang,J.,
Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collamore,A.,
Cooke,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Fato,S., Ferreira,P., Fitzhugh,W., Fitzhugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I.,
Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Larroque,K.,
Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Marquis,N.,
Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Mldrim,J.,
Meneus,L., Mlhova,T., Mlhova,T., Mlhova,T., Mlhova,V.,
Mphuy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H.,
O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
Pollara,V., Pollara,V., Roman,J., Rosetti,M., Riley,R., Rise,C.,
Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S.,
Schnupack,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Straus,N., Subramanian,A., Talamas,J., Testaye,S.,
Theodore,J., Topham,K., Travers,M., Travis,N., Triggillo,J.,
Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
TITLE
Direct Submission
JOURNAL
Submitted (14-JUN-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE
4 (bases 1 to 191434)
AUTHORS
Birten,B., Bastien,V., Bloom,T., Boguslavsky,L., Boukhgalter,B.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhgalter,B.,
Cammarata,J., Chang,J., Chazaro,B., Choepel,Y., Collamore,A.,
Cooke,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Fato,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J.,
Gardyna,S., Ginde,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Mldrim,J., Meneus,L., Mlhova,T., Mlhova,V.,
Mphuy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schnupack,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
TITLE
Direct Submission
JOURNAL
Submitted (16-JUN-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT
On Jul 16, 2002 this sequence version replaced gi:20043143.

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All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIR

Web site: <http://www.seq.wi.mit.edu>

Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)

----- Project Information

Center project name: L11935

Center clone name: 522\_B\_15

# FEATURES

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      20896..21016
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      21079..21098
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      21150..21200
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                complement(23784..24062)
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                24327..24350
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                /rpt_family="AluSc"
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                complement(30791..31103)
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 Best local similarity 90.9%; Pred. No. 85;  
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAGACTTCTGACTACATCA 22  
 DB 137915 AAGACTTCTGATTACATCAA 137936

RESULT 9  
 AL772303/c 197437 bp DNA linear HTG 13-JUL-2002  
 LOCUS Mus musculus chromosome 2 clone RP23-185P20, \*\*\* SEQUENCING IN  
 DEFINITION  
 ACCESSION AL772303  
 VERSION AL772303.3 GI:21748303  
 KEYWORDS HTG; HTGS; PHASE1; HTGS\_ACTIVEPIN; HTGS\_DRAFT; HTGS\_FULLTOP.  
 SOURCE house mouse.  
 ORGANISM Mus musculus;

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 REFERENCE 1 Submitted (12-JUL-2002) Wellcome Trust Sanger Institute, Hinxton,  
 Cambridgeshire, CB10 1SA, UK. E-mail enquiries:  
 humbrey@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk  
 On Jul 14, 2002 this sequence version replaced gi:21540206.  
 COMMENT ----- Genome Center  
 Center: Wellcome Trust Sanger Institute  
 Center code: SC  
 Web site: <http://www.sanger.ac.uk>

Contact: humquery@esanger.ac.uk  
 ----- Project Information  
 Center project name: BM185P20  
 ----- Summary Statistics  
 Assembly program: XGAP4; version 4.5  
 Chemistry: Dye-terminator; 100% of reads  
 Consensus quality: 196057 bases at least Q40  
 Consensus quality: 196481 bases at least Q30  
 Consensus quality: 196768 bases at least Q20  
 Insert size: 196937; sum-of-contigs  
 Quality coverage: 5.88x in Q20 bases; sum-of-contigs quality  
 coverage: 5.99x in Q20 bases; agarose-1p  
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 \* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 6 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.  
 \* 1 46842: contig of 46842 bp in length  
 \* 46843 46942: gap of 100 bp  
 \* 46943 51200: contig of 4258 bp in length  
 \* 51301 51300: gap of 100 bp  
 \* 51301 62621: contig of 11321 bp in length  
 \* 62622 62721: gap of 100 bp  
 \* 62722 92981: contig of 30260 bp in length  
 \* 92982 93081: gap of 100 bp  
 \* 93082 143175: contig of 50094 bp in length  
 \* 143176 143275: gap of 100 bp  
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 93082. 143175  
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## FEATURES

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 59973 a 40264 c 38730 g 57570 t 500 others  
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 ORIGIN  
 Query Match 85.5%; Score 18.6; DB 2; Length 197437;  
 Best Local Similarity 90.9%; Pred. No. 85;  
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 AAGACTTGTGAGTAACATCAA 22  
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 Db 189362 AAGACTTCTAATTAACATCAA 189341

RESULT 10 AC127239 223761 bp DNA linear HTG 17-JUL-2002  
 AC127239  
 LOCUS  
 DEFINITION Mus musculus chromosome UNK clone RP24-381A23, WORKING DRAFT

SEQUENCE, 7 unordered pieces.  
 AC127239  
 AC127239.1 GI:21747652  
 HTG: HTGS\_PHAISEL; HTGS\_DRAFT.  
 KEYWORDS  
 SOURCE  
 ORGANISM  
 Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 1 (bases 1 to 223761)  
 McPherson, J.D. and Waterston, R.H.  
 The sequence of Mus musculus clone  
 Unpublished  
 2 (bases 1 to 223761)  
 McPherson, J.D. and Waterston, R.H.  
 Direct Submission  
 Submitted (14-JUL-2002) Genome Sequencing Center, 4444 Forest Park  
 Parkway, St. Louis, MO 63108, USA  
 3 (bases 1 to 223761)  
 McPherson, J.D. and Waterston, R.H.  
 Direct Submission  
 Submitted (17-JUL-2002) Genome Sequencing Center, 4444 Forest Park  
 Parkway, St. Louis, MO 63108, USA  
 COMMENT  
 ----- Genome Center -----  
 Center: Washington University Genome Sequencing Center  
 Center code: WUGSC  
 Web site: <http://genome.wustl.edu/gsc/index.shtml>  
 Contact: [submissions@watson.wustl.edu](mailto:submissions@watson.wustl.edu)  
 ----- Project Information -----  
 Center project name: M\_BB0381A23  
 ----- Summary Statistics -----  
 Sequencing vector: M13; 0%  
 Sequencing vector: Plasmid; 100%  
 Chemistry: Dye-terminator Big Dye; 100% of reads  
 Assembly program: Phrap; version 0.990319  
 Consensus quality: 221262 bases at least Q40  
 Consensus quality: 221869 bases at least Q30  
 Consensus quality: 222233 bases at least Q20  
 -----  
 \* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 7 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.  
 \* 1 6570: contig of 6570 bp in length  
 \* 6571 6670: gap of unknown length  
 \* 6671 22737: contig of 16067 bp in length  
 \* 22738 22837: gap of unknown length  
 \* 22838 48321: contig of 25484 bp in length  
 \* 48322 48421: gap of unknown length  
 \* 48422 118533: contig of 70112 bp in length  
 \* 118534 118633: gap of unknown length  
 \* 118634 212339: contig of 93706 bp in length  
 \* 212340 212439: gap of unknown length  
 \* 212440 216233: contig of 3794 bp in length  
 \* 216234 216333: gap of unknown length  
 \* 216334 223761: contig of 7428 bp in length.  
 Location/Qualifiers  
 1. 223761  
 /organism="Mus musculus"  
 /db\_xref="taxon:10090"  
 /chromosome="UNK"  
 /clone="RP24-381A23"  
 1. 6570  
 /note="assembly\_name:Contig10"  
 6671. 22737  
 /note="assembly\_name:Contig11"  
 22838. 48321

## FEATURES

## source

misc\_feature  
 /note="assembly\_name:Contig10"  
 6671. 22737  
 /note="assembly\_name:Contig11"  
 22838. 48321  
 misc\_feature



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misc_feature      /note="assembly_name:Contig12"
                  48422..118533
misc_feature      /note="assembly_name:Contig13"
                  118634..212339
misc_feature      /note="assembly_name:Contig14"
                  212440..216233
misc_feature      /note="assembly_name:Contig18"
                  216334..223761
misc_feature      /note="assembly_name:Contig19"
                  71420 a 41598 c 40871 g 69263 t 609 others
BASE COUNT      71420 a 41598 c 40871 g 69263 t
ORIGIN

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Query Match      85.5%; Score 18.8; DB 2; Length 223761;
Best Local Similarity 90.9%; Pred. No. 84;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

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QY      1 AAGACTTCTGACTACATCA 22
        ||||||| ||| |||||||
DB      87767 AAGACTTCTGACTACATCA 87788

```

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RESULT 11
AC098888      229247 bp      DNA      linear      ROD 21-JUN-2002
DEFINITION    Mus musculus clone RP23-122N8, complete sequence.
ACCESSION     AC098888
VERSION       AC098888.4 GI:19909498
KEYWORDS      HTG.
SOURCE        house mouse.
ORGANISM      Mus musculus
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

```

```

REFERENCE     1 (bases 1 to 229247)
AUTHORS      McPherson,J.D. and Waterston,R.H.
TITLE        The sequence of Mus musculus clone
JOURNAL      Unpublished
REFERENCE     2 (bases 1 to 229247)
AUTHORS      McPherson,J.D. and Waterston,R.H.
TITLE        Direct Submission
JOURNAL      Submitted (05-NOV-2001) Genome Sequencing Center, 4444 Forest Park
              Parkway, St. Louis, MO 63108, USA

```

```

REFERENCE     3 (bases 1 to 229247)
AUTHORS      McPherson,J.D. and Waterston,R.H.
TITLE        Direct Submission
JOURNAL      Submitted (03-APR-2002) Genome Sequencing Center, 4444 Forest Park
              Parkway, St. Louis, MO 63108, USA
REFERENCE     4 (bases 1 to 229247)
AUTHORS      McPherson,J.D. and Waterston,R.H.
TITLE        Direct Submission
JOURNAL      Submitted (21-JUN-2002) Genome Sequencing Center, 4444 Forest Park
              Parkway, St. Louis, MO 63108, USA

```

On Apr 3, 2002 this sequence version replaced gi:17105321.

```

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
Contact: submissions@watson.wustl.edu
Project Information
Center project name: M_BA0122N08
-----
FEATURES
source
1..229247
location/qualifiers
/organism="Mus musculus"
/db_xref="taxon:10090"
/clone="RP23-122N8"

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BASE COUNT      70714 a 42415 c 42708 g 73410 t
ORIGIN
Query Match      85.5%; Score 18.8; DB 10; Length 229247;
Best Local Similarity 90.9%; Pred. No. 83;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

```

QY      1 AAGACTTCTGACTACATCA 22
        ||||||| ||| |||||||
DB      83202 AAGACTTCTGACTACATCA 83223

```

```

RESULT 12
AC099415      317515 bp      DNA      linear      HTG 05-JUN-2002
LOCUS         Mus musculus chromosome UNK clone RP23-122D8, WORKING DRAFT
DEFINITION    AC099415
ACCESSION     AC099415
VERSION       AC099415.3 GI:21326409
KEYWORDS      HTG: HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEFIN.
SOURCE        house mouse.
ORGANISM      Mus musculus
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

```

```

REFERENCE     1 (bases 1 to 317515)
AUTHORS      McPherson,J.D. and Waterston,R.H.
TITLE        The sequence of Mus musculus clone
JOURNAL      Unpublished
REFERENCE     2 (bases 1 to 317515)
AUTHORS      McPherson,J.D. and Waterston,R.H.
TITLE        Direct Submission
JOURNAL      Submitted (14-NOV-2001) Genome Sequencing Center, 4444 Forest Park
              Parkway, St. Louis, MO 63108, USA
REFERENCE     3 (bases 1 to 317515)
AUTHORS      McPherson,J.D. and Waterston,R.H.
TITLE        Direct Submission
JOURNAL      Submitted (05-JUN-2002) Genome Sequencing Center, 4444 Forest Park
              Parkway, St. Louis, MO 63108, USA

```

On Jun 5, 2002 this sequence version replaced gi:20069750.

```

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
Contact: submissions@watson.wustl.edu
Project Information
Center project name: M_BA0122D08
-----

```

```

----- Summary Statistics -----
Sequencing vector: M13; 16%
Sequencing vector: plasmid; 84%
Chemistry: Dye-primer EM; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 293243 bases at least Q40
Consensus quality: 303369 bases at least Q30
Consensus quality: 309642 bases at least Q20
Insert size: 224000; agarose-fp
Insert size: 322378; sum-of-ctrls
Quality coverage: 37.21 in Q20 bases; agarose-fp
Quality coverage: 24.94 in Q20 bases; sum-of-ctrls
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 56 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 1211: contig of 1211 bp in length
* 1212 1311: gap of unknown length
* 1312 2357: contig of 1046 bp in length
* 2358 2457: gap of unknown length
* 2458 3574: contig of 1117 bp in length
* 3574 3674: gap of unknown length
* 3675 4817: contig of 1143 bp in length
* 4818 4917: gap of unknown length
* 4918 6015: contig of 1098 bp in length
* 6016 6016: gap of unknown length

```

```

* 6116 7259: contig of 1144 bp in length
* 7260 7359: gap of unknown length
* 7360 8831: contig of 1472 bp in length
* 8832 8931: gap of unknown length
* 8932 10179: contig of 1248 bp in length
* 10180 10279: gap of unknown length
* 10280 11586: contig of 1307 bp in length
* 11587 11686: gap of unknown length
* 11687 12940: contig of 1254 bp in length
* 12941 13040: gap of unknown length
* 13041 14154: contig of 1114 bp in length
* 14155 14254: gap of unknown length
* 14255 15728: contig of 1474 bp in length
* 15729 15828: gap of unknown length
* 15829 17212: contig of 1384 bp in length
* 17213 17312: gap of unknown length
* 17313 18620: contig of 1308 bp in length
* 18621 18720: gap of unknown length
* 18721 20283: contig of 1563 bp in length
* 20284 20384: gap of unknown length
* 20384 21676: contig of 1293 bp in length
* 21677 21776: gap of unknown length
* 21777 23147: contig of 1371 bp in length
* 23148 23247: gap of unknown length
* 23248 24781: contig of 1534 bp in length
* 24782 24882: gap of unknown length
* 24882 26627: contig of 1746 bp in length
* 26628 26727: gap of unknown length
* 26728 28451: contig of 1724 bp in length
* 28452 28551: gap of unknown length
* 28552 29863: contig of 1312 bp in length
* 29864 29963: gap of unknown length
* 29964 31941: contig of 1978 bp in length
* 31942 32041: gap of unknown length
* 32042 33660: contig of 1619 bp in length
* 33661 33760: gap of unknown length
* 33761 34889: contig of 1129 bp in length
* 34890 36451: gap of unknown length
* 36452 36551: gap of unknown length
* 36552 38157: contig of 1606 bp in length
* 38158 38257: gap of unknown length
* 38258 40052: contig of 1795 bp in length
* 40053 40152: gap of unknown length
* 40153 41725: contig of 1573 bp in length
* 41726 41825: gap of unknown length
* 41826 42996: contig of 1171 bp in length
* 42997 43096: gap of unknown length
* 43097 44639: contig of 1543 bp in length
* 44640 44739: gap of unknown length
* 44740 46047: contig of 1308 bp in length
* 46048 46147: gap of unknown length
* 46148 47831: contig of 1684 bp in length
* 47832 47931: gap of unknown length
* 47932 49690: contig of 1759 bp in length
* 49691 49790: gap of unknown length
* 49791 51094: contig of 1304 bp in length
* 51095 51194: gap of unknown length
* 51195 53501: contig of 2307 bp in length
* 53502 53601: gap of unknown length
* 53602 55272: contig of 1671 bp in length
* 55273 55372: gap of unknown length
* 55373 57428: contig of 2056 bp in length
* 57429 57528: gap of unknown length
* 57529 59819: contig of 2291 bp in length
* 59820 59919: gap of unknown length
* 59920 61717: contig of 1798 bp in length
* 61718 61817: gap of unknown length
* 61818 64311: contig of 2494 bp in length
* 64312 67042: contig of 2631 bp in length
* 67043 67142: gap of unknown length
* 67143 69238: contig of 2096 bp in length

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```

* 69239 69338: gap of unknown length
* 69339 71520: contig of 2182 bp in length
* 71521 71620: gap of unknown length
* 71621 74319: contig of 2699 bp in length
* 74320 74419: gap of unknown length
* 74420 76137: contig of 1718 bp in length
* 76138 76237: gap of unknown length
* 76238 79611: contig of 3374 bp in length
* 79612 82702: contig of 2991 bp in length
* 82703 82802: gap of unknown length
* 82803 84448: contig of 1646 bp in length
* 84449 84548: gap of unknown length
* 84549 87522: contig of 2874 bp in length
* 87523 89088: contig of 1566 bp in length
* 89089 89188: gap of unknown length
* 89189 92189: contig of 3001 bp in length
* 92190 92289: gap of unknown length
* 92290 96604: contig of 4315 bp in length
* 96605 96705: gap of unknown length
* 96706 98834: contig of 2130 bp in length
* 98835 98934: gap of unknown length
* 98935 109620: contig of 10686 bp in length
* 109621 109720: gap of unknown length
* 109721 177250: contig of 67530 bp in length
* 177251 177350: gap of unknown length
* 177351 317515: contig of 140165 bp in length.

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## FEATURES

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misc_feature      1..1211
                  /organism="Mus musculus"
                  /db_xref="taxon:10090"
                  /chromosome="1"
                  /clone="RP23-122D8"
misc_feature      1312..2357
                  /note="assembly_name:Contig189"
misc_feature      2458..3574
                  /note="assembly_name:Contig193"
misc_feature      3675..4817
                  /note="assembly_name:Contig141"
misc_feature      4916..6015
                  /note="assembly_name:Contig142"
misc_feature      6116..7259
                  /note="assembly_name:Contig147"
misc_feature      7360..8831
                  /note="assembly_name:Contig159"
misc_feature      8932..10179
                  /note="assembly_name:Contig163"
misc_feature      10280..11586
                  /note="assembly_name:Contig165"
misc_feature      11687..12940
                  /note="assembly_name:Contig170"
misc_feature      13041..14154
                  /note="assembly_name:Contig174"
misc_feature      14255..15728
                  /note="assembly_name:Contig177"
misc_feature      15829..17212
                  /note="assembly_name:Contig179"
misc_feature      17313..18620
                  /note="assembly_name:Contig180"

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## Query Match

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Best Local Similarity 85.5%; Score 18.8; DB 2; Length 317515;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

QY 1 AGACTTCGAGTACATCA 22

DB 17367 AGACTTCGAGTACATCA 17388

RESULT 13  
AE011043/c

LOCUS	AE011043	10243 bp	DNA	linear	BCT 03-APR-2002
DEFINITION	Methanosarcina acetivorans str. C2A, section 388 of 534 of the complete genome.				
ACCESSION	AE011043 AE010239				
VERSION	AE011043.1 GI:19917413				
KEYWORDS					
SOURCE	Methanosarcina acetivorans C2A.				
ORGANISM	Methanosarcina acetivorans C2A				
REFERENCE	Archaea: Euryarchaeota: Methanococci: Methanosarcinales; Methanosarcinaceae: Methanosarcina.				
AUTHORS	1 (bases 1 to 10243) Galagan, J.E., Nussbaum, C., Roy, A., Endlitz, M.G., Macdonald, P., Fitzhugh, W., Calvo, S., Engels, R., Smirnov, S., Atnoor, D., Brown, A., Allen, N., Naylor, J., Stange-Thomann, N., DeVellano, K., Johnson, R., Linton, L., McEwan, P., McKernan, K., Talamas, J., Tirrell, A., Ye, W., Zimmer, A., Barber, R.D., Cann, I., Graham, D.E., Grahame, D.A., Krzycki, J.A., Leigh, J.A., Li, W., Liu, J., Mukhopadhyay, B., Reeve, J.N., Smith, K., Springer, T.A., Umayam, L.A., White, O., White, R.H., de Macario, E.C., Ferry, J.G., Jarrell, K.F., Jing, H., Macario, A.J.L., Paulsen, I., Pritchett, M., Sowers, K.R., Swanson, R.V., Zinder, S.H., Lander, E., Metcalf, W.W. and Birren, B.				
TITLE	The Genome of <i>M. acetivorans</i> Reveals Extensive Metabolic and Physiological Diversity				
JOURNAL	Genome Res. 12 (4), 532-542 (2002)				
MEDLINE	21929760				
PUBMED	11932238				
REFERENCE	2 (bases 1 to 10243)				
AUTHORS	Birren, B.				
TITLE	Direct Submission				
JOURNAL	Submitted (20-MAR-2002) Center for Genome Research, Whitehead Institute, Nine Cambridge Center, Cambridge, MA 02141, USA				
FEATURES	Location/Qualifiers				
Source	1..10243				
gene	/organism="Methanosarcina acetivorans C2A"				
CDS	/strain="C2A"				
gene	/db_xref="taxon:188937"				
CDS	complement(88..603)				
gene	/gene="MA3371"				
CDS	complement(88..603)				
gene	/gene="MA3371"				
CDS	/codon_start=1				
gene	/transl_table=11				
CDS	/product="predicted protein"				
gene	/protein_id="AA06740.1"				
CDS	/db_xref="GI:19917414"				
gene	/translation="MIRENNIMAIMGREGRGPLDIDIGIVVSGIGSGNSQNRNOF NGLFSGIDGEGLYIGPSNPFORTISIGIGVIGSGNGLAKLGHFRAGLQML PPIVYVGVSGASFLALPVBEGKAAATIPILNVPAAVYQENIKELRQGTETLML LRSYISGRTP"				
gene	complement(859..1488)				
CDS	/gene="MA3372"				
gene	complement(859..1488)				
CDS	/gene="MA3372"				
gene	/codon_start=1				
gene	/transl_table=11				
CDS	/product="conserved hypothetical protein"				
gene	/protein_id="AA06741.1"				
CDS	/db_xref="GI:19917415"				
gene	/translation="MKIAVENENQOQSIFEPFTAVVEEDGVKKVILNRENQVCA ARGMAAVMAVGAIDIKQIDGVKVVASIEGICAGSCTFOAAGPIFIYDVSVDLDTI KGMLEETEKROEPRKFDTEPLEPREENGDSINIEDIMEFPPDLSKTLIPYLK NGFERNLDVIGCGHMKFVTDLGMGEFETVNESTNRKTRVRAQT"				
gene	1981..3642				
CDS	/gene="11VD"				
gene	/note="MA3373"				
CDS	1981..3642				
gene	/gene="11VD"				
CDS	/codon_start=1				
gene	/transl_table=11				
CDS	/product="dihydroxy-acid dehydratase"				
gene	/protein_id="AA06742.1"				
CDS	/db_xref="GI:19917416"				
gene	/translation="MRSDIIEGPERAPNRSILKATGVTSEMKKPTIAVNSNDII PGHMLNKLAEAVKACIRNAGVPEFHHVGVCDGAMGHEMKSLSRITIEDTIE LAYKAVDGMVLPSCDITVPCGHMAARLDIPATVVTGCMLEGYDDNDRIDTISV SEGVAFVSAGKISIEPEFKLENI.SCGASGSCGAPANTAMCMTALSLSPGCTAH AVDAKKILIAKESGERIVELVKNLTPKIVTQKSENAINVDAVGSSTVTLPLA LAHERGLPLEAFDELRSKTPHLISLRGPNFLIHPDRAGVAVVQKSLKHLID QLVNKGRTGLENDELEIVNPLNMEITPLENPPIHAEGIAVLKSGAPGSAVKOA ADPKNHVYTPAKVYDCEDEAMKSLIADGVKPGDIPVIRYRGPGKSGEMLAATA ALGRLLSVALVTDGREGSGTGRGCGIGVSPKSDGPTAIYVDSGLIETINPERA LNKISEELKOKASFPKPTKITGYIARVRAVHSAVNTGIVD"				
gene	complement(3805..4149)				
CDS	/gene="MA3374"				
gene	complement(3805..4149)				
CDS	/gene="MA3374"				
gene	/codon_start=1				
gene	/transl_table=11				
CDS	/product="conserved hypothetical protein"				
gene	/protein_id="AA06743.1"				
CDS	/db_xref="GI:19917417"				
gene	/translation="MNIADVGRKKFNCISFGSROTKQLAMEKKEIGEYMTDH MNAVEFETIHTOSKAEYVEGNGILRHFLARLRKTKYKSLKEMKYSVLL MKHRNKELSTF"				
gene	complement(4053..4487)				
CDS	/gene="MA3375"				
gene	complement(4053..4487)				
CDS	/gene="MA3375"				
gene	/codon_start=1				
gene	/transl_table=11				
CDS	/product="conserved hypothetical protein"				
gene	/protein_id="AA06744.1"				
CDS	/db_xref="GI:19917418"				
gene	/translation="MNCPRCKSSNHTKNGIVGQKQKCHDCGYNSVELSTAPSL VAKQALQVLESLGRLGRLGVSHSVQKMKFQCEIEELKSENESISIVELDEMH TYIGKKNIAGSGLLIELGKNSSTALLVAERKLDVSGKN"				
gene	4675..4950				
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gene	4675..4950				
CDS	/gene="MA3376"				
gene	/codon_start=1				
gene	/transl_table=11				
CDS	/product="predicted protein"				
gene	/protein_id="AA06745.1"				
CDS	/db_xref="GI:19917419"				
gene	/translation="MGVIATRLFGGLIKRSQSDVEFTGKIGYPAVKDIINFA TKSYSVDLIDLNGIPETIYNTDEVETREIERSORTREYRPEY"				
gene	5699..7393				
CDS	/gene="MA3377"				
gene	5699..7393				
CDS	/gene="MA3377"				
gene	/codon_start=1				
gene	/transl_table=11				
CDS	/product="conserved hypothetical protein"				
gene	/protein_id="AA06746.1"				
CDS	/db_xref="GI:19917420"				
gene	/translation="MCTTLITTRGASKDSMTVAHSDDELGNORITVYPADHEGA MROYFEHYRPRYLVTYERGNPERPAHHNSQSPNPPDLTGPIGTIPVRRHYAFD RYVGLNENHNLMEECTNGAKIOPAHVASAEAKNSKHRLTFYSQLSVALERCKA REAVLGLKLDKTYISTEETLLVADEDAWFEKCALPDEQYHSAMVAQVQDEY FVAANERIRIQIEDNRDII.CSELKGLKGLNLMWPEIDPLMLSSVANGENHPYU SLRWVAVLVRVNPDLALSPWVTRGFVDGSGYAIIDYPSIKPGLILOVFPALYR DHYEGTQFDLIRGVAAAGPYGDPNRPFGYQNNVTNENHWGAGERPISFYQGYT YVNLKRTPEIIPASGICLPGPDVSYTTCPPAPAAQELPERIOSDPIPOVQPT AWMAFDFVSNMARNYORMTRVILPQQKITEVEVQKTILOWDECCKRSQLEARDL LTRUGKENANDEVKQMHKLGHALLIARTSDGICMLPNQBPONIGYSSKWLALTNKDG PTSTEMKP"				
gene	complement(7624..7914)				
CDS	/gene="MA3378"				
gene	complement(7624..7914)				
CDS	/gene="MA3378"				
gene	/codon_start=1				
gene	/transl_table=11				
CDS	/product="predicted protein"				
gene	/protein_id="AA06747.1"				

gene  
CDS  
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/translation="MISMDSMEKIFGKTAQMTVLKNIENONEPYVLSGIAETGLSH  
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complement(8740..10059)  
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/transl\_table=1  
/product="amino acid transporter"  
/protein\_id="AA06748.1"  
/db\_xref="GI:19917422"  
/translation="MEEVKNSGNYGKRESEELKRDLSLEITVIGDILGAGIVYM  
GKAAGLAGNMWAFLEFAGFTASLSALSYELSSMTPRGAETVPARKAFGERIGLTV  
GLVIVYVAITASAVALGFRYSNIFGGIILGALSILSLVWVYIKSARLAI  
LMTFELSGMLITVIGIPYIGVNYEPFSLGVEASLIIFAFGEDIVRLSQE  
TKDAEKTTPKRLILSIFETIYICVAVASVLDPOVLGISEIPLAEVAANAFGKA  
FVLSWLTALBSTMTVIVVMIGSRIVYGMANGSLPKILARVHOKLTPMTAICGIA  
FFSLFVFLGDIATVANIATNPMTIVTFYINISLILKRYTPERKRPFPVPSIGRFP  
IFPALGALSAVFLPSQIGKEVMTLIGFELAGISALVILKTRKENEL"

BASE COUNT 3037 a 2258 c 2111 g 2837 t

ORIGIN

Query Match 83.6%; Score 18.4; DB 1; Length 10243;  
Best Local Similarity 95.0%; Pred. No. 1.8e+02;  
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 AAGACTTCTAGTACATC 20  
1 |||||  
Db 2740 AGGACTTCTAGTACATC 2721

RESULT 14  
AE010860/c 10467 bp DNA linear BCT 03-APR-2002  
LOCUS  
DEFINITION Methanosarcina acetivorans str. C2A, section 205 of 534 of the  
complete genome.  
ACCESSION AE010860 AE010299  
VERSION AE010860.1 GI:19915697  
KEYWORDS  
SOURCE Methanosarcina acetivorans C2A.  
ORGANISM Methanosarcina acetivorans C2A.  
REFERENCE  
AUTHORS Galagan, J.E., Nusbbaum, C., Roy, A., Endrizzi, M.G., MacDonald, P.,  
Ritzhugh, W., Calvo, S., Engels, R., Smirnov, S., Alnour, D., Brown, A.,  
Allen, N., Naylor, J., Stange-Thomann, N., Dearrellano, K., Johnson, R.,  
Linton, L., McEwan, P., McKernan, K., Talamas, J., Tittell, A., Ye, W.,  
Zimmer, A., Barber, R.D., Cann, I., Graham, D.E., Grahame, D.A.,  
Guss, A., Hedderich, R., Ingram-Smith, C., Kuetner, C.H.,  
Krzyski, J.A., Leigh, J.A., Li, W., Liu, J., Mukhopadhyay, B.,  
Reeve, J.N., Smith, K., Springer, T.A., Umayam, L.A., White, O.,  
White, R.H., de Macario, E.C., Perry, J.G., Jarrell, K.F., Jing, H.,  
Macario, A.J.L., Paulsen, I., Pritchett, M., Swens, K.R.,  
Swanson, R.V., Zinder, S.H., Lander, E., Metcalf, W.W. and Birren, B.  
The genome of *M. acetivorans* Reveals Extensive Metabolic and  
Physiological Diversity  
Genome Res. 12 (4), 532-542 (2002)

JOURNAL  
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AUTHORS Birren, B.  
TITLE Direct Submission  
JOURNAL Submitted (20-MAR-2002) Center for Genome Research, Whitehead  
Institute, Nine Cambridge Center, Cambridge, MA 02141, USA  
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Tue Mar 18 16:16:18 2003

us-09-836-439-6.rge

Page 18

Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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GenCore version 5.1.4.p5.4578  
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OM nucleic - nucleic search, using sw model

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 PT Nucleic acids encoding human immune/hematopoietic antigen polypeptides,  
 PT useful for preventing, diagnosing and/or treating cancers and  
 PT metastasis -  
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 PS Disclosure; SEQ ID NO 37150; 3071pp + Sequence Listing; English.  
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 XX AAk54951 to AAk64702 encode the human immune/hematopoietic antigen (I)  
 CC amino acid sequences given in AAk62170 to AAk91921. (I) have cytostatic  
 CC activity, and can be used in gene therapy and vaccine production. (I)  
 CC and polynucleotides may be used in the prevention, diagnosis and  
 CC treatment of diseases associated with inappropriate (I) expression. For  
 CC example, they may be used to treat disorders associated with decreased  
 CC expression by rectifying mutations or deletions in a patient's genome  
 CC that affect the activity of (I) by expressing inactive proteins or to



CC supplement the patients own production of (I). Additionally, (I)  
CC polynucleotides may be used to produce the secreted (I) by inserting  
CC the nucleic acids into a host cell and culturing the cell to express the  
CC protein. (I) proteins and polynucleotides may be used to prevent,  
CC diagnose and treat immune/hematopoietic-related diseases, especially  
CC to AK6/894 represent human immune/hematopoietic antigen genomic  
CC sequences from the present invention. AK54942 to AK54950 and AKM82169  
CC represent sequences used in the exemplification of the present invention.  
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PR 18-JUN-1999; 99US-0139461.  
PR 18-JUN-1999; 99US-0139462.  
PR 18-JUN-1999; 99US-0139463.  
PR 18-JUN-1999; 99US-0139750.  
PR 21-JUN-1999; 99US-0139753.  
PR 22-JUN-1999; 99US-0139817.  
PR 23-JUN-1999; 99US-0139899.  
PR 23-JUN-1999; 99US-0140354.  
PR 24-JUN-1999; 99US-0140695.  
PR 28-JUN-1999; 99US-0140823.  
PR 29-JUN-1999; 99US-0140991.  
PR 30-JUN-1999; 99US-0141287.  
PR 01-JUL-1999; 99US-0141842.  
PR 02-JUL-1999; 99US-0142154.  
PR 06-JUL-1999; 99US-0142055.  
PR 08-JUL-1999; 99US-0142390.  
PR 09-JUL-1999; 99US-0142803.  
PR 12-JUL-1999; 99US-0142977.  
PR 13-JUL-1999; 99US-0143542.  
PR 14-JUL-1999; 99US-0143624.  
PR 15-JUL-1999; 99US-0144005.  
PR 16-JUL-1999; 99US-0144085.  
PR 16-JUL-1999; 99US-0144086.  
PR 19-JUL-1999; 99US-0144325.  
PR 19-JUL-1999; 99US-0144331.  
PR 19-JUL-1999; 99US-0144332.  
PR 19-JUL-1999; 99US-0144333.  
PR 19-JUL-1999; 99US-0144334.  
PR 19-JUL-1999; 99US-0144335.  
PR 20-JUL-1999; 99US-0144352.  
PR 20-JUL-1999; 99US-0144632.  
PR 21-JUL-1999; 99US-0144884.  
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PR 21-JUL-1999; 99US-0145086.  
PR 22-JUL-1999; 99US-0145088.  
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PR 22-JUL-1999; 99US-0145088.  
PR 22-JUL-1999; 99US-0145089.  
PR 22-JUL-1999; 99US-0145192.  
PR 23-JUL-1999; 99US-0145145.  
PR 23-JUL-1999; 99US-0145218.  
PR 23-JUL-1999; 99US-0145224.  
PR 26-JUL-1999; 99US-0145276.  
PR 27-JUL-1999; 99US-0145913.  
PR 27-JUL-1999; 99US-0145918.  
PR 27-JUL-1999; 99US-0145919.  
PR 28-JUL-1999; 99US-0145951.  
PR 02-AUG-1999; 99US-0146386.

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PR 02-AUG-1999; 99US-0146388.
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PR 13-AUG-1999; 99US-0148664.
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PR 18-AUG-1999; 99US-0149426.
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PR 23-AUG-1999; 99US-0149902.
PR 23-AUG-1999; 99US-0149930.
PR 25-AUG-1999; 99US-0150566.
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PR 27-AUG-1999; 99US-0151066.
PR 27-AUG-1999; 99US-0151080.
PR 30-AUG-1999; 99US-0151303.
PR 31-AUG-1999; 99US-0151438.
PR 01-SEP-1999; 99US-0151930.
PR 07-SEP-1999; 99US-0152363.
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PR 24-SEP-1999; 99US-0156458.
PR 28-SEP-1999; 99US-0156596.
PR 29-SEP-1999; 99US-0157117.
PR 04-OCT-1999; 99US-0157753.
PR 05-OCT-1999; 99US-0157865.
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PR 07-OCT-1999; 99US-0158232.
PR 08-OCT-1999; 99US-0158369.
PR 12-OCT-1999; 99US-0159293.
PR 13-OCT-1999; 99US-0159294.
PR 13-OCT-1999; 99US-0159295.
PR 14-OCT-1999; 99US-0159329.
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PR 14-OCT-1999; 99US-0159637.
PR 14-OCT-1999; 99US-0159638.
PR 18-OCT-1999; 99US-0159584.
PR 21-OCT-1999; 99US-0160741.
PR 21-OCT-1999; 99US-0160767.
PR 21-OCT-1999; 99US-0160768.
PR 21-OCT-1999; 99US-0160770.
PR 21-OCT-1999; 99US-0160814.
PR 21-OCT-1999; 99US-0160815.
PR 22-OCT-1999; 99US-0160860.
PR 22-OCT-1999; 99US-0160981.
PR 22-OCT-1999; 99US-0160989.
PR 25-OCT-1999; 99US-0161404.
PR 25-OCT-1999; 99US-0161405.
PR 25-OCT-1999; 99US-0161406.
PR 26-OCT-1999; 99US-0161359.
PR 26-OCT-1999; 99US-0161360.
PR 26-OCT-1999; 99US-0161361.

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PR 28-OCT-1999; 99US-0161920.
PR 28-OCT-1999; 99US-0161992.
PR 28-OCT-1999; 99US-0161993.
PR 29-OCT-1999; 99US-0162142.

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Query Match 80.9%; Score 17.8; DB 21; Length 801;
Best Local Similarity 90.5%; Pred. No. 56;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 AAGACTTCTGAGTACATCA 21
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DB 543 AAGACTTCTGAGTACATCA 563

```

```

RESULT 3
AA003999/c
ID AA003999 standard; DNA; 877 BP.

```

```

XX AA003999;

```

```

DT 03-SEP-1990 (first entry)

```

```

DE Sequence complementary to dystrophin gene.

```

```

XX X-chromosome; ornithine transcarbamylase deficiency;
KW muscular dystrophy; dystrophin; ds.

```

```

XX Synthetic.

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```

PN EP364255-A.

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XX 18-APR-1990.

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XX 11-OCT-1989; 89EP-0310424.

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XX 12-OCT-1988; 88US-0256689.

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PA (BAYU ) BAYLOR UNIV COLLEGE.

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PI Caskay CT, Chamberlain JS, Gibbs RAL, Rainer JE, Nguyen PN;

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DR WPI; 1990-117752/16.

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PT Multiplex genomic DNA amplification for deletion detection -
  useful for detecting X-linked diseases such as ornithine
  transcarbamylase deficiency and X-linked muscular dystrophy.

```

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XX Claim 9; Page 19; 32pp; English.

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```

CC Paired oligonucleotide primers are used in detecting deletions
  CC specifically of the X and Y chromosomes.

```

```

CC Dystrophin gene may be isolated this way.

```

```

XX Sequence 877 BP; 289 A; 160 C; 131 G; 297 T; 0 other;

```

```

Query Match 80.9%; Score 17.8; DB 11; Length 877;
Best Local Similarity 90.5%; Pred. No. 56;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 AAGACTTCTGAGTACATCA 21
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DB 132 AAGCTTCTGAGTACATCA 112

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RESULT 4
AAC50782
ID AAC50782 standard; DNA; 1353 BP.

```

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XX AAC50782;

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AC AAC50782;

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```

DT 18-OCT-2000 (first entry)
XX Arabidopsis thaliana DNA fragment SEQ ID NO: 66116.
DE

```

XX Hybridisation assay; genetic mapping; gene expression control;  
KW protein identification; signal transduction pathway;  
KW metabolic pathway; promoter; termination sequence; ss.  
OS *Arabidopsis thaliana*.  
PN EP1033405-A2.  
XX  
PD 06-SEP-2000.  
XX  
PF 25-FEB-2000; 2000EP-0301439.  
XX  
PR 25-FEB-1999; 99US-0121825.  
PR 05-MAR-1999; 99US-0123180.  
PR 09-MAR-1999; 99US-0123548.  
PR 23-MAR-1999; 99US-0125788.  
PR 25-MAR-1999; 99US-0126264.  
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PR 01-APR-1999; 99US-0127462.  
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PR 30-APR-1999; 99US-0132407.  
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PR 18-MAY-1999; 99US-0134768.  
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PR 29-JUN-1999; 99US-0140991.  
PR 30-JUN-1999; 99US-0141287.  
PR 01-JUL-1999; 99US-0141842.  
PR 01-JUL-1999; 99US-0142154.  
PR 02-JUL-1999; 99US-0142055.  
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PR 08-JUL-1999; 99US-0142803.  
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PR 19-JUL-1999; 99US-0144325.  
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PR 19-JUL-1999; 99US-0144333.  
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PR 21-JUL-1999; 99US-0145086.  
PR 21-JUL-1999; 99US-0145088.  
PR 22-JUL-1999; 99US-0145085.  
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PR 22-JUL-1999; 99US-0145089.  
PR 22-JUL-1999; 99US-0145192.  
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PR 27-JUL-1999; 99US-0145918.  
PR 27-JUL-1999; 99US-0145919.  
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PR 23-AUG-1999; 99US-0149902.  
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PR 25-AUG-1999; 99US-0150566.  
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PR 27-AUG-1999; 99US-0151065.  
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PR 27-AUG-1999; 99US-0151080.  
PR 30-AUG-1999; 99US-0151303.

PR 31-AUG-1999; 99US-0151438.  
 PR 01-SEP-1999; 99US-0151930.  
 PR 07-SEP-1999; 99US-0152363.  
 PR 10-SEP-1999; 99US-0153070.  
 PR 13-SEP-1999; 99US-0153758.  
 PR 15-SEP-1999; 99US-0154018.  
 PR 16-SEP-1999; 99US-0154039.  
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 PR 06-OCT-1999; 99US-0157865.  
 PR 07-OCT-1999; 99US-0158029.  
 PR 08-OCT-1999; 99US-0158232.  
 PR 12-OCT-1999; 99US-0158369.  
 PR 13-OCT-1999; 99US-0159293.  
 PR 13-OCT-1999; 99US-0159294.  
 PR 13-OCT-1999; 99US-0159295.  
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 PR 22-OCT-1999; 99US-0160989.  
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 PR 26-OCT-1999; 99US-0161360.  
 PR 26-OCT-1999; 99US-0161361.  
 PR 28-OCT-1999; 99US-0161920.  
 PR 28-OCT-1999; 99US-0161992.  
 PR 28-OCT-1999; 99US-0161993.  
 PR 29-OCT-1999; 99US-0162142.

Query Match 80.9%; Score 17.8; DB 21; Length 1353;  
 Best Local Similarity 90.5%; Pred. No. 59;  
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 AAGACTTCTGACTTAACATCA 21  
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 Db 543 AAGACTTCTGACTTAACATCA 563

## RESULT 5

AAK54111/c  
 ID AAK54111 standard; cDNA; 513 BP.

XX AAK54111;

XX 16-NOV-2001 (first entry)

DE Murine transport and binding associated protein encoding cDNA SEQ ID 676.

XX Murine; liver; gene library; amino acid synthesis; binding protein;

KW cell metabolism; energy metabolism; fatty acid metabolism; synthesis;

KW phospholipid metabolism; purine; pyrimidine; nucleoside; nucleotide;

KW replication; transcription; translation; transport protein; ss.

XX Mus musculus.

XX DE20103510-U1.  
 XX 07-JUN-2001.  
 XX 28-FEB-2001; 2001DE-2003510.  
 XX 02-DEC-1999; 99DE-1058160.  
 XX (LION-) LION BIOSCIENCE AG.  
 XX WPI; 2001-368570/39.  
 XX Gene library containing sequences with specific 3'-ends and no polyA  
 PT tail, encoding proteins involved in a wide range of cellular processes  
 PT  
 PS  
 XX  
 CC This invention describes a novel gene library (A) comprises a gene  
 CC sequence (or its part) encoding a protein involved in amino acid  
 CC synthesis, cellular/energy metabolism, metabolism of  
 CC fatty acids/phospholipids, synthesis or breakdown of  
 CC purines/pyrimidines/nucleosides/nucleotides, DNA  
 CC replication/transcription/translation, or is a transport/binding protein.  
 CC (A) are produced that correspond to the 3'-end of mRNA but without the  
 CC polyA tail. They can be prepared more efficiently and with less effort  
 CC than conventional libraries. AAK53436-AAK54275 represent fragments of the  
 CC gene library described in the method of the invention.  
 XX  
 SQ Sequence 513 BP; 143 A; 125 C; 107 G; 137 T; 1 other;

Query Match 78.2%; Score 17.2; DB 22; Length 513;  
 Best Local Similarity 86.4%; Pred. No. 1e+02;  
 Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 AAGACTTCTGACTTAACATCA 22  
 ||||| ||||| ||||| |||||  
 Db 22 AAGACTTCTGACTTAACATCA 1

RESULT 6

ABL07736  
 ID ABL07736 standard; cDNA; 7626 BP.

XX ABL07736;

DT 26-MAR-2002 (first entry)

XX Drosophila melanogaster expressed polynucleotide SEQ ID NO 17690.

DE Drosophila; developmental biology; cell signalling; insecticide;

KW pharmaceutical; gene; ss.

XX Drosophila melanogaster.

XX WO200171042-A2.

XX 27-SEP-2001.

XX 23-MAR-2001; 2001WO-US09231.

XX 23-MAR-2000; 2000US-191637P.

XX 11-JUL-2000; 2000US-0614150.

XX (PEKE ) PE CORP NY.

XX Venter JC, Adams M, Li PWD, Myers EW;

XX WPI; 2001-656860/75.

XX P-PSDB; ABB63633.

XX New isolated nucleic acid detection reagent for detecting 1000 or more

PT genes from *Drosophila* and for elucidating cell signalling and cell-cell interactions.

XX Claim 1: SEQ ID NO 17690: 21bp + Sequence Listing; English.

PS

CC The invention relates to an isolated nucleic acid detection reagent capable of detecting 1000 or more genes from *Drosophila*. The invention is useful in developmental biology and in elucidating cell signalling and cell-cell interactions in higher eukaryotes for the development of insecticides, therapeutics and pharmaceutical drugs. The invention discloses genomic DNA sequences (AB101840-AB16175) and the encoded DNA sequences (AB857737-AB872072).

CC The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp://wipo.int/pub/published\_pct\_sequences.

CC

CC Sequence 7626 BP; 1991 A; 1673 C; 1791 G; 2171 T; 0 other;

SO

Query Match 78.2%; Score 17.2; DB 23; Length 7626;  
Best Local Similarity 86.4%; Pred. No. 1.3e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AAGACTTCTGAGTACATCA 22  
|||||  
DB 5862 AAGACTTCTGAGTACATCA 5883

RESULT 7  
AAH68529 standard; DNA; 349980 BP.

XX

AC AAH68529;

XX

DT 26-SEP-2001 (first entry)

XX

DE C glutamicum coding sequence fragment SEQ ID NO: 7064.

XX

KW Coryneform bacterium; amino acid synthesis; vitamin; saccharide;

KM organic acid synthesis; ds.

XX

OS Corynebacterium glutamicum.

XX

PN EP1108790-A2.

XX

PD 20-JUN-2001.

XX

PF 18-DEC-2000; 2000EP-0127668.

XX

PR 16-DEC-1999; 99JP-0377484.

PR 07-APR-2000; 2000JP-0159162.

PR 03-AUG-2000; 2000JP-0280988.

XX

PA (KYOWA) KYOWA HAKKO KOGYO KK.

PI Nakagawa S, Mizoguchi H, Ando S, Hayashi M, Ochiai K, Yokoi H;  
Tateishi N, Senoh A, Ikeda M, Ozaki A;

PI

DR WPI: 2001-376931/40.

XX

XX Novel polynucleotides derived from Coryneform bacteria, for identifying mutation point of a gene, measuring expression of a gene, analysing expression profile or pattern of a gene and identifying homologous gene

PT

XX

PS Disclosure: SEQ ID NO: 7064; 246pp + Sequence Listing; English.

XX

CC The present invention provides a number of nucleotide and protein sequences from the Coryneform bacterium *Corynebacterium glutamicum*. These are useful for identifying the mutation point of a gene derived from a mutant of *Corynebacterium glutamicum*, measuring expression amount and analysing the expression profile or expression pattern of a gene derived from *Corynebacterium glutamicum*, and identifying a homologue of a gene derived

CC from *Coryneform* bacterium. *Coryneform* bacteria are useful for producing amino acids, nucleic acids, vitamins, saccharides and organic acids, particularly L-lysine. The present sequence is a nucleic acid described in the exemplification of the invention.

CC Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from the European Patent Office.

CC

CC Sequence 349980 BP; 82466 A; 95954 C; 90516 G; 81044 T; 0 other;

SO

Query Match 78.2%; Score 17.2; DB 22; Length 349980;  
Best Local Similarity 86.4%; Pred. No. 1.8e+02;  
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AAGACTTCTGAGTACATCA 22  
|||||  
DB 232860 AAGACTTCTGAGTACATCA 232881

RESULT 8  
AAC57234 standard; DNA; 461 BP.

XX

AC AAC57234;

XX

DT 25-JAN-2001 (first entry)

XX

DE *Pinus radiata* transcription factor DNA sequence #631.

XX

KW Plant; transcription factor; gene expression; eucalyptus; pine; acacia; poplar; sweetgum; teak; mahogany; bZIP; G-box binding factor;

KM basic helix-loop-helix zipper; homeotic; homeodomain; homeobox; MADS; homeodomain zipper; LIM domain; AP2; ERBs; zinc finger domain;

KW type 2 Cys2His2; CCAAT box element; MYB; ss.

XX

OS *Pinus radiata*.

XX

PN WO200053724-A2.

XX

PD 14-SEP-2000.

XX

PF 09-MAR-2000; 2000WO-US06112.

XX

PR 11-MAR-1999; 99US-0266513.

PR 18-AUG-1999; 99US-0149485.

XX

PA (GENE-) GENESIS RES & DEV CORP LTD.  
(FLEET-) FLEETCHER CHALLENGE FORESTS LTD.

PI

PI Wood M, McGrath A, Shenk MA, Glenn M;

PI

DR WPI: 2000-579369/54.

XX

XX New isolated polynucleotide encoding a plant transcription factor for producing a plant e.g. a woody plant, preferably eucalyptus or pine, having modified gene expression or modified activity of a polypeptide

PT

XX

PS Claim 1: Page 624; 747pp; English.

XX

CC The present invention relates to novel plant transcription factors from *Eucalyptus grandis* or *Pinus radiata*. The present sequence is the coding sequence for one such transcription factor. The transcription factor may be used to produce a plant having modified gene expression such as a woody plant e.g. a eucalyptus, pine, acacia, poplar, sweetgum, teak, or mahogany species or to modify the activity of a polypeptide in a plant. The transcription factors of the present invention are members from the following families of regulatory proteins: bZIP, bZIP family of G-box binding factors, basic helix-loop-helix zipper, LIM domain, AP2 homeotic/homeodomain/homeobox/MADS, homeodomain zipper, LIM domain, AP2 and ERBs, zinc finger domains of type 2 Cys2His2, CCAAT box elements and MYB.

Sequence 461 BP; 163 A; 97 C; 94 G; 107 T; 0 other;  
Query Match 76.4%; Score 16.8; DB 21; Length 461;  
Best Local Similarity 90.0%; Pred. No. 1.6e+02;  
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
QY 3 GACTCTGAGTACATCAA 22  
||||| ||||| |||||  
Db 408 GACTTCTCAGTACAGTCAA 427  
RESULT 9  
AAK66626  
ID AAK66626 standard; DNA; 21477 BP.  
AC AAK66626;  
XX  
XX 06-NOV-2001 (first entry)  
DT  
XX  
XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:21438.  
DE  
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;  
KW cytostatic; gene therapy; vaccine; metastasis; ds.  
XX  
XX Homo sapiens.  
OS  
PN WO200157182-A2.  
XX  
XX 09-AUG-2001.  
PD  
XX  
XX 17-JAN-2001; 2001WO-US01354.  
PF  
XX  
XX 31-JAN-2000; 2000US-0179065.  
PR 04-FEB-2000; 2000US-0180628.  
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PR 15-MAR-2000; 2000US-0189874.  
PR 17-MAR-2000; 2000US-0190076.  
PR 18-APR-2000; 2000US-0198123.  
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PR 07-JUN-2000; 2000US-0209467.  
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PR 26-JUL-2000; 2000US-0220964.  
PR 14-AUG-2000; 2000US-0224518.  
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PR 14-AUG-2000; 2000US-0225213.  
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PR 14-AUG-2000; 2000US-0225266.  
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PR 14-AUG-2000; 2000US-0225447.  
PR 14-AUG-2000; 2000US-0225757.  
PR 14-AUG-2000; 2000US-0225758.  
PR 14-AUG-2000; 2000US-0225759.  
PR 18-AUG-2000; 2000US-0226279.  
PR 22-AUG-2000; 2000US-0226681.  
PR 22-AUG-2000; 2000US-0226868.  
PR 22-AUG-2000; 2000US-0227182.  
PR 23-AUG-2000; 2000US-0227009.  
PR 30-AUG-2000; 2000US-0228924.  
PR 01-SEP-2000; 2000US-0229287.  
PR 01-SEP-2000; 2000US-0229343.  
PR 01-SEP-2000; 2000US-0229344.  
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PR 05-SEP-2000; 2000US-0229513.  
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PR 08-SEP-2000; 2000US-0232080.  
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PR 12-SEP-2000; 2000US-0231968.  
PR 14-SEP-2000; 2000US-0232397.  
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PR 14-SEP-2000; 2000US-0233064.  
PR 14-SEP-2000; 2000US-0233065.  
PR 21-SEP-2000; 2000US-0234223.  
PR 21-SEP-2000; 2000US-0234274.  
PR 25-SEP-2000; 2000US-0234997.  
PR 25-SEP-2000; 2000US-0234998.  
PR 26-SEP-2000; 2000US-0235484.  
PR 27-SEP-2000; 2000US-0235834.  
PR 27-SEP-2000; 2000US-0235836.  
PR 29-SEP-2000; 2000US-0236327.  
PR 29-SEP-2000; 2000US-0236367.  
PR 29-SEP-2000; 2000US-0236368.  
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PR 29-SEP-2000; 2000US-0236370.  
PR 02-OCT-2000; 2000US-0236802.  
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PR 21-SEP-2000; 2000US-0234274.  
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 PR 17-NOV-2000; 2000US-0249297.  
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 PR 05-DEC-2000; 2000US-0251988.  
 PR 05-DEC-2000; 2000US-0251988.  
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 PR 08-DEC-2000; 2000US-0251989.  
 PR 08-DEC-2000; 2000US-0251990.

PR 11-DEC-2000; 2000US-0254097.  
 PR 05-JAN-2001; 2001US-0259678.  
 PA (HUMA-) HUMAN GENOME SCI INC.  
 PI Rosen CA, Barash SC, Ruben SM;  
 XX WPI; 2001-483426/52.  
 DR  
 XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,  
 PT useful for preventing, diagnosing and/or treating cancers and  
 PT metastasis -  
 PS  
 XX  
 XX Disclosure; SEQ ID NO 21437; 3071bp + Sequence Listing; English.  
 CC  
 CC AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)  
 CC amino acid sequences given in AAK82170 to AAK91921. (I) have cytosolic  
 CC activity, and can be used in gene therapy and vaccine production. (I)  
 CC proteins and polynucleotides may be used in the prevention, diagnosis and  
 CC treatment of diseases associated with inappropriate (I) expression. For  
 CC example, they may be used to treat disorders associated with decreased  
 CC expression by rectifying mutations or deletions in a patient's genome  
 CC that affect the activity of (I) by expressing inactive proteins or to  
 CC supplement the patient's own production of (I). Additionally, (I)  
 CC polynucleotides may be used to produce the secreted (I), by inserting the  
 CC nucleic acids into a host cell and culturing the cell to express the  
 CC protein. (I) proteins and polynucleotides may be used to prevent,  
 CC diagnose and treat immune/hematopoietic-related diseases, especially  
 CC cancers and cancer metastases of hematopoietic-derived cells. AAK64703  
 CC to AAK87694 represent human immune/hematopoietic antigen genomic  
 CC sequences from the present invention. AAK54942 to AAK54950 and AAK62169  
 CC represent sequences used in the exemplification of the present invention.  
 CC  
 XX Sequence 21480 BP; 5315 A; 5005 C; 5257 G; 5903 T; 0 other;  
 SQ  
 Query Match 76.4%; Score 16.8; DB 22; Length 21480;  
 Best Local Similarity 90.0%; Pred. No. 2.3e+02;  
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 Db 7222 AGCCTTCGATGATCAATCA 7241  
 QY 2 AGACTTCGATGATCAATCA 21  
 II |||||  
 DB 7222 AGCCTTCGATGATCAATCA 7241  
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 AAS01375/C  
 ID AAS01375 standard; CDNA; 3114 BP.  
 XX  
 AC AAS01375;  
 XX  
 DT 04-JUL-2001 (first entry)  
 XX  
 XX Human TANGO 405 cDNA sequence.  
 DE  
 XX  
 XX Human; TANGO 210; clone jthla152h06; TANGO 364; TANGO 366; dectin-2;  
 KW INTERCEPT 394; INTERCEPT 400; TANGO 405; cellular process regulator;  
 KW gene therapy; growth modulator; proliferation; cell differentiation;  
 KW lymphocyte; bone marrow cell migration; leukaemia; lymphoma;  
 KW autoimmune disorder; ss.  
 KW  
 XX  
 OS Homo sapiens.  
 XX  
 OS  
 FH Key Location/Qualifiers  
 FT CDS 154..783  
 FT /tag= a  
 FT /product= "TANGO 405 protein"  
 FT /note= "The ORF is specifically claimed"  
 FT sig\_peptide 154..297  
 FT /tag= b  
 FT mat\_peptide 298..780  
 FT /tag= c  
 XX  
 PN WC200118016-A1.



XX 15-MAR-2001.  
PD 30-JUN-2000; 2000MO-US18174.  
XX 10-SEP-1999; 99US-0393996.  
PR (MILL-) MILLENNIUM PHARM INC.  
XX  
PI Fraser CC, Sharp JD, Wighton N, Myers PS, Goodearl ADJ;  
XX WPI: 2001-183280/18.  
DR P-PSDB; AAU00479.  
XX  
PT Isolated nucleic acid molecules encoding proteins useful as modulating  
PT agents in regulating a variety of cellular processes are used for  
XX treating e.g. cancer and autoimmune disorders -  
PS Claim 2; Fig 6A-6C; 326pp; English.  
XX  
CC The present sequence encoding for human TANGO 405 protein is isolated  
CC from cDNA clone jh1a152h06 from a human mixed lymphocyte reaction cDNA  
CC library. It is 1 of 6 novel human proteins which include TANGO 210  
CC (AAU00469), TANGO 364 (AAU00471), TANGO 366 (AAU00472), INTERCEPT 394  
CC (AAU00473), and INTERCEPT 400 (AAU00476). Novel sequences for murine  
CC TANGO 210 (AAU00470), INTERCEPT 400 (AAU00477), TANGO 405 (AAU00480) and  
CC a rat INTERCEPT 400 (AAU00478) sequence are also described. The nucleic  
CC acids encoding these novel proteins are useful as modulating agents in  
CC regulating a variety of cellular processes and can be used to express  
CC the proteins in a host cell in gene therapy applications. Human and  
CC murine TANGO 405 proteins show sequence homology to murine decin-2.  
CC TANGO 405 modulates growth, proliferation, survival, differentiation,  
CC activity, morphology and movement/migration of human lymphocytes and  
CC bone marrow cells and tissues and can be used to prevent, diagnose or  
CC treat leukemia, lymphomas and autoimmune disorders.  
XX  
SQ Sequence 3114 BP; 1001 A; 527 C; 517 G; 1069 T; 0 other;  
Query Match 74.5%; Score 16.4; DB 22; Length 3114;  
Best Local Similarity 94.4%; Pred. No. 3e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
OY 5 CTTCTGAGTAACATCA 22  
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DB 1885 CTTCTGAGTAACATCA 1868  
RESULT 12  
ID ABA18026  
AC ABA18026 standard; DNA; 29329 BP.  
XX  
AC ABA18026;  
XX  
DT 23-JAN-2002 (first entry)  
XX  
DE Human nervous system related polynucleotide SEQ ID NO 10357.  
XX  
KW Human; nootropic; neuroprotective; cytoskeletal; dermatological; virologic;  
KW immunosuppressive; antiinflammatory; anti-HIV; antibacterial; vulnery;  
KW antiparkinsonian; antisticking; antianaemic; antiarthritic; cancer;  
KW antineumatic; hepatotoxic; cerebroprotective; antiinflammatory;  
KW antiallergic; antidiabetic; antilucer; anticonvulsant; antifungal;  
KW antiparasitic; cardiac; immune disorder; cardiovascular disorder;  
KW neurological disease; infection; nephrotoxic; gene therapy; vaccine; ds.  
XX  
OS Homo sapiens.  
XX  
PN WO200159063-A2.  
XX  
PD 16-AUG-2001.  
XX  
PF 17-JAN-2001; 2001MO-US01334.  
XX

PR 31-JAN-2000; 2000US-0179065.  
PR 04-FEB-2000; 2000US-0180628.  
PR 24-FEB-2000; 2000US-0184664.  
PR 02-MAR-2000; 2000US-0186350.  
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PR 17-MAR-2000; 2000US-0190076.  
PR 18-APR-2000; 2000US-0198123.  
PR 19-MAY-2000; 2000US-0205515.  
PR 07-JUN-2000; 2000US-0209467.  
PR 28-JUN-2000; 2000US-0214886.  
PR 30-JUN-2000; 2000US-0215135.  
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PR 18-AUG-2000; 2000US-0225759.  
PR 22-AUG-2000; 2000US-0226279.  
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PR 01-SEP-2000; 2000US-0228927.  
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PR 05-SEP-2000; 2000US-0229345.  
PR 05-SEP-2000; 2000US-0229509.  
PR 06-SEP-2000; 2000US-0229513.  
PR 06-SEP-2000; 2000US-0230437.  
PR 08-SEP-2000; 2000US-0230438.  
PR 08-SEP-2000; 2000US-0231242.  
PR 08-SEP-2000; 2000US-0231243.  
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PR 08-SEP-2000; 2000US-0232080.  
PR 12-SEP-2000; 2000US-0232081.  
PR 14-SEP-2000; 2000US-0231968.  
PR 14-SEP-2000; 2000US-0232397.  
PR 14-SEP-2000; 2000US-0232398.  
PR 14-SEP-2000; 2000US-0232400.  
PR 14-SEP-2000; 2000US-0232401.  
PR 14-SEP-2000; 2000US-0233063.  
PR 14-SEP-2000; 2000US-0233064.  
PR 21-SEP-2000; 2000US-0233065.  
PR 21-SEP-2000; 2000US-0234223.  
PR 21-SEP-2000; 2000US-0234274.  
PR 25-SEP-2000; 2000US-0234597.  
PR 25-SEP-2000; 2000US-0234598.  
PR 26-SEP-2000; 2000US-0234998.  
PR 27-SEP-2000; 2000US-0235834.  
PR 27-SEP-2000; 2000US-0235836.  
PR 29-SEP-2000; 2000US-0236327.  
PR 29-SEP-2000; 2000US-0236367.  
PR 29-SEP-2000; 2000US-0236368.  
PR 29-SEP-2000; 2000US-0236369.  
PR 29-SEP-2000; 2000US-0236570.  
PR 02-OCT-2000; 2000US-0236802.

PS	Disclosure; SEQ ID NO 10357; 1701bp + Sequence Listing; English.
XX	
CC	The invention relates to novel genes (ABA11004-ABA21534) and proteins
CC	(ABA14678-ABA18001) useful for preventing, treating or ameliorating
CC	medical conditions e.g. by protein or gene therapy. The genes are
CC	isolated from a range of human tissues disclosed in the specification.
CC	The nucleic acids, proteins, antibodies and (ant)agonists are useful
CC	in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast
CC	and ovarian cancer and other cancers of the adrenal gland, bone, bone
CC	marrow, breast, gastrointestinal tract, liver, lung, or urogenital;
CC	(b) immune disorders e.g. Addison's disease, diabetes mellitus, Crohn's
CC	hemolytic anaemia, autoimmune thyroiditis, diabetes mellitus and ulcerative
CC	disease, multiple sclerosis, rheumatoid arthritis and ulcerative
CC	colitis; (c) cardiovascular disorders such as myocardial ischaemia;
CC	(d) wound healing; (e) neurological diseases e.g. cerebral anoxia and
CC	epilepsy; and (f) infectious diseases such as viral, bacterial, fungal
CC	and parasitic infections.
CC	Note: The sequence data for this patent did not form part of the
CC	printed specification, but was obtained in electronic format directly
CC	from WIPO at ftp.wipo.int/pub/published_pat_sequences.
XX	
SO	Sequence 29329 BP; 6864 A; 6756 C; 7225 G; 8484 T; 0 other;
QY	
Query Match	74.5%; Score 16.4; DB 22; Length 29329;
Best Local Similarity	94.4%; Pred. No. 3.6e+02;
Matches 1/; Conservative	0; Mismatches 1; Indels 0; Gaps 0;
DB	
3 GACTTCTGAGTAAACAATC 20	
8110 GACTTCTGAGAAACAATC 8127	
RESULT 13	
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ID	ABA20511 standard; DNA: 29329 BP.
XX	
AC	ABA20511;
XX	
DT	23-JAN-2002 (first entry)
XX	
DE	
XX	Human nervous system related polynucleotide SEQ ID NO 12842.
XX	
XX	Human; nootropic; neuroprotective; cytostatic; dermatological; virucide;
KW	immunosuppressive; anti-inflammation; anti-HIV; antibacterial; vulnery;
KW	antiparkinsonia; antisickling; antianaemic; antiarthritic; cancer;
KW	antihepatic; hepatotropic; cerebroprotective; antiinflammatory;
KW	antiallergic; antidiabetic; antitumor; anticonvulsant; antifungal;
KW	antiparasitic; cardiac; immune disorder; cardiovascular disorder;
XX	neurological disease; infection; nephrotropic; gene therapy; vaccine; ds.
XX	
OS	Homo sapiens.
XX	
PN	WO200159063-A2.
XX	
PD	16-AUG-2001.
XX	
PF	17-JAN-2001; 2001WO-US01334.
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PR	31-JAN-2000; 2000US-0179065.
PR	04-FEB-2000; 2000US-0180628.
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PR	02-MAR-2000; 2000US-0186350.
PR	16-MAR-2000; 2000US-0189874.
PR	17-MAR-2000; 2000US-0190076.
PR	18-APR-2000; 2000US-0198123.
PR	19-MAY-2000; 2000US-0205515.
PR	07-JUN-2000; 2000US-0209467.
PR	28-JUN-2000; 2000US-0214986.
PR	30-JUN-2000; 2000US-0215135.
PR	07-JUL-2000; 2000US-0216647.
PR	07-JUL-2000; 2000US-0216880.
PR	11-JUL-2000; 2000US-0217487.



CC colitis; (c) cardiovascular disorders such as myocardial ischaemia;  
CC (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and  
CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal  
CC and parasitic infections.  
CC Note: The sequence data for this patent did not form part of the  
CC printed specification, but was obtained in electronic format directly  
CC from WIPO at ftp.wipo.int/pub/published\_pcl\_sequences.  
XX  
SQ Sequence 29329 BP; 6864 A; 6756 C; 7225 G; 8484 T; 0 other;  
  
Query Match 74.5%; Score 16.4; DB 22; Length 29329;  
Best Local Similarity 94.4%; Pred. No. 3.6e+02;  
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
  
QY 3 GACTCTGTGATACATC 20  
DB 8110 GACTCTGTGATACATC 8127  
  
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XX  
AC AAK70791;  
XX  
DT 06-NOV-2001 (first entry)  
XX  
DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:25603.  
XX  
KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;  
KW cytostatic; gene therapy; vaccine; metastasis; ds.  
XX  
OS Homo sapiens.  
XX  
PN WO200157182-A2.  
XX  
PD 09-AUG-2001.  
XX  
PF 17-JAN-2001; 2001WO-US01354.  
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PR 31-JAN-2000; 2000US-0179065.  
PR 04-FEB-2000; 2000US-0180628.  
PR 24-FEB-2000; 2000US-0184664.  
PR 02-MAR-2000; 2000US-0186350.  
PR 16-MAR-2000; 2000US-0189874.  
PR 17-MAR-2000; 2000US-0190076.  
PR 18-APR-2000; 2000US-0198123.  
PR 19-MAY-2000; 2000US-0205515.  
PR 07-JUN-2000; 2000US-0209467.  
PR 28-JUN-2000; 2000US-0214886.  
PR 30-JUN-2000; 2000US-0215135.  
PR 07-JUL-2000; 2000US-0216647.  
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PR 29-SEP-2000; 2000US-0236327.  
PR 29-SEP-2000; 2000US-0236367.  
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PR 29-SEP-2000; 2000US-0236369.  
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PR 29-SEP-2000; 2000US-0236802.  
PR 02-OCT-2000; 2000US-0237037.  
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PR 02-OCT-2000; 2000US-0237039.  
PR 02-OCT-2000; 2000US-0237040.  
PR 13-OCT-2000; 2000US-0239937.  
PR 13-OCT-2000; 2000US-0240960.  
PR 20-OCT-2000; 2000US-0241221.  
PR 20-OCT-2000; 2000US-0241785.  
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PR 17-NOV-2000; 2000US-0249207.  
PR 17-NOV-2000; 2000US-0249208.

PR 17-NOV-2000; 2000US-0249209.  
PR 17-NOV-2000; 2000US-0249210.  
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PR 17-NOV-2000; 2000US-0249216.  
PR 17-NOV-2000; 2000US-0249217.  
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PR 17-NOV-2000; 2000US-0249244.  
PR 17-NOV-2000; 2000US-0249245.  
PR 17-NOV-2000; 2000US-0249264.  
PR 17-NOV-2000; 2000US-0249265.  
PR 17-NOV-2000; 2000US-0249297.  
PR 17-NOV-2000; 2000US-0249299.  
PR 01-DEC-2000; 2000US-0249300.  
PR 01-DEC-2000; 2000US-0250160.  
PR 05-DEC-2000; 2000US-0250391.  
PR 05-DEC-2000; 2000US-0251030.  
PR 05-DEC-2000; 2000US-0251988.  
PR 06-DEC-2000; 2000US-0256719.  
PR 08-DEC-2000; 2000US-0251856.  
PR 08-DEC-2000; 2000US-0251868.  
PR 08-DEC-2000; 2000US-0251869.  
PR 08-DEC-2000; 2000US-0251989.  
PR 11-DEC-2000; 2000US-0251990.  
PR 05-JAN-2001; 2001US-0259678.  
XX  
XX  
XX (HUMA-) HUMAN GENOME SCI INC.  
XX  
XX Rosen CA, Barash SC, Ruben SM;  
XX  
XX WPI; 2001-483426/52.  
XX  
XX  
XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,  
XX useful for preventing, diagnosing and/or treating cancers and  
XX metastasis.

Disclosure; SEQ ID NO 25603; 3071pp + Sequence Listing: English.

CC AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)  
CC amino acid sequences given in AAK82170 to AAK91921. (I) have cytosolic  
CC activity, and can be used in gene therapy and vaccine production. (I)  
CC proteins and polynucleotides may be used in the prevention, diagnosis and  
CC treatment of diseases associated with inappropriate (I) expression. For  
CC example, they may be used to treat disorders associated with decreased  
CC expression by rectifying mutations or deletions in a patient's genome  
CC that affect the activity of (I) by expressing inactive proteins or to  
CC supplement the patient's own production of (I). Additionally, (I)  
CC polynucleotides may be used to produce the secreted (I), by inserting  
CC the nucleic acids into a host cell and culturing the cell to express the  
CC protein. (I) proteins and polynucleotides may be used to prevent,  
CC diagnose and treat immune/hematopoietic-related diseases, especially  
CC to AAK87694 represent human immune/hematopoietic-derived cells. AAK64703  
CC sequences from the present invention. AAK54942 to AAK54950 and AAK82169  
CC represent sequences used in the exemplification of the present invention.  
XX  
XX  
XX Sequence 29329 BP; 6864 A; 6756 C; 7225 G; 8484 T; 0 other;

Query Match 74.5%; Score 16.4; DB 22; Length 29329;  
Best Local Similarity 94.4%; Pred. No. 3.6e+02;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 GACTTGTGATTAACATC 20  
DB 8110 GACTTGTGAGAAACATC 8127  
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RESULT 15

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AC  
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XX 07-NOV-2001 (first entry)  
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XX  
XX Human; immune; hematopoietic; immune/hematopoietic antigen; cancer;  
XX cytostatic; gene therapy; vaccine; metastasis; ds.  
XX  
XX Homo sapiens.  
XX  
XX WO200157182-A2.  
XX  
XX 09-AUG-2001.  
XX  
XX 17-JAN-2001; 2001WO-US01354.  
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XX 31-JAN-2000; 2000US-0179065.  
XX 04-FEB-2000; 2000US-0180628.  
XX 24-FEB-2000; 2000US-0184664.  
XX 02-MAR-2000; 2000US-0186350.  
XX 16-MAR-2000; 2000US-0189874.  
XX 17-MAR-2000; 2000US-0190076.  
XX 18-APR-2000; 2000US-0198123.  
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XX 07-JUN-2000; 2000US-0209467.  
XX 28-JUN-2000; 2000US-0214886.  
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XX 11-JUL-2000; 2000US-0217487.  
XX 14-JUL-2000; 2000US-0217496.  
XX 26-JUL-2000; 2000US-0218290.  
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XX 14-AUG-2000; 2000US-0224518.  
XX 14-AUG-2000; 2000US-0224519.  
XX 14-AUG-2000; 2000US-0225213.  
XX 14-AUG-2000; 2000US-0225214.  
XX 14-AUG-2000; 2000US-0225266.  
XX 14-AUG-2000; 2000US-0225267.  
XX 14-AUG-2000; 2000US-0225268.  
XX 14-AUG-2000; 2000US-0225270.  
XX 14-AUG-2000; 2000US-0225447.  
XX 14-AUG-2000; 2000US-0225757.  
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XX 18-AUG-2000; 2000US-0226279.  
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XX 22-AUG-2000; 2000US-0226688.  
XX 23-AUG-2000; 2000US-0227182.  
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XX      05-JAN-2001; 2001US-0259678.  
  
IPA      (HUMA-) HUMAN GENOME SCI INC.  
XX  
XX      Rosen CA, Barash SC, Ruben SM;  
DR       WPI; 2001-483426/52.  
  
PT       Nucleic acids encoding human immune/hematopoietic antigen polypeptides,  
PP        useful for preventing, diagnosing and/or treating cancers and  
PS         metastasis -  
XX  
PS        Disclosure; SEQ ID NO 33324; 3071pp + Sequence Listing; English.  
CC      AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)  
CC amino acid sequences given in AA#M82170 to AA#G91921. (I) have cytostatic  
CC activity, and can be used in gene therapy and vaccine production. (I)  
CC proteins and polynucleotides may be used in the prevention, diagnosis and  
CC treatment of diseases associated with inappropriate (I) expression. For  
CC example, they may be used to treat disorders associate with decreased  
CC expression by rectifying mutations or deletions in a patient's genome  
CC to supplement the patients own production of (I). Additionally, (I)  
CC polynucleotides may be used to produce the secreted (I), by inserting the  
CC nucleic acids into a host cell and culturing the cell to express the  
CC protein. (II) proteins and polynucleotides may be used to prevent,  
CC diagnose and treat immune/haematopoeitic-related diseases, especially  
CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703  
CC to AAK87694 represent human immune/haematopoietic antigen genomic  
CC sequences from the present invention. AAK54942 to AAK54950 and AA#M82169  
CC represent sequences used in the exemplification of the present invention.  
XX  
XX      Sequence 29329 BP; 6864 A; 6756 C; 7225 G; 8484 T; 0 other;
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GenCore version 5.1.4-p5\_4578  
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OM nucleic - nucleic search, using sw model

Run on: March 17, 2003, 10:23:12 ; Search time 777.688 Seconds  
(without alignments)  
458.154 Million cell updates/sec

Title: US-09-836-439-6

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Scoring table: IDENTITY\_NUC  
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Searched: 16154066 seqs, 8097743376 residues  
Total number of hits satisfying chosen parameters: 32308132

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Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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C 4	17.8	80.9	418	14	B0622226 fc11c.pk0
C 5	17.8	80.9	509	12	B0882650 sae94g07.
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C 12	17.4	79.1	549	10	AW738913 g016b06.y
C 13	17.4	79.1	608	17	AQ525332 HS_5226.B
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C 15	17.4	79.1	692	13	BJ173371 BJ173371
C 16	17.4	79.1	734	13	BJ167340 BJ167340
C 17	17.2	78.2	106	10	BE339863 EST343923
C 18	17.2	78.2	185	17	AZ242233
C 19	17.2	78.2	236	10	BB528073
C 20	17.2	78.2	258	10	BB301743
C 21	17.2	78.2	325	17	AZ746982
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C 23	17.2	78.2	390	12	BF451198
C 24	17.2	78.2	391	12	BF333029
C 25	17.2	78.2	396	17	AZ104881
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C 29	17.2	78.2	425	9	AT663816
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C 31	17.2	78.2	436	9	AU014847
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C 33	17.2	78.2	454	17	AQ769890
C 34	17.2	78.2	460	10	BE579476
C 35	17.2	78.2	473	10	AM220604
C 36	17.2	78.2	482	9	AT267625
C 37	17.2	78.2	482	10	AM398852
C 38	17.2	78.2	487	10	BE462943
C 39	17.2	78.2	489	9	AU080451
C 40	17.2	78.2	489	9	AA270973
C 41	17.2	78.2	492	12	BF116785
C 42	17.2	78.2	511	12	BE016374
C 43	17.2	78.2	512	10	BE689588
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C 45	17.2	78.2	518	17	AQ416223

# ALIGNMENTS

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ACCESSION AA970603  
VERSION AA970603.1 GI:3145110  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 347)  
AUTHORS NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.  
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
Tumor Gene Index  
JOURNAL Unpublished (1997)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgaps-remail.nih.gov  
This clone is available royalty-free through LNL ; contact the  
IMAGE Consortium (info@image.llnl.gov) for further information.  
Insert Length: 1053 Std Error: 0.00  
Seq primer: -40ml3 fwd. ET from Amersham  
High quality sequence stop: 344.  
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/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:1579337"

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/clone_11b="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NBHL19W, Testis NHT, and B-cell
NCI-CGAP-GCB1) were mixed, and ss circles were made in
vitro. Following HAP purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from pools of 5,000 clones made
from the same 3 libraries. The pools consisted of
I.M.A.G.E. clones 297480-302087, 682632-687239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo."
BASE COUNT      91 a      75 c      63 g      118 t
ORIGIN

Query Match      83.6%; Score 18.4; DB 9; Length 347;
Best Local Similarity 95.0%; Pred. No. 4.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY      3 GACTTCTGAGTACCAATCAA 22
Db      335 GACTTCTGAGTACCAATCAA 316

RESULT 2
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LOCUS      od6e07.s1 NCI-CGAP-GCB1 Homo sapiens CDNA clone IMAGE:1369092,
DEFINITION      mRNA sequence.
ACCESSION      AA837289
VERSION      AA837289.1 GI:2912488
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 395)
REFERENCE      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
TITLE      Tumor Gene Index
JOURNAL      Unpublished (1997)
COMMENT      Contact: Robert Strausberg, Ph.D.
Email: cgabs-remail.nih.gov
Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
Ph.D., Gerald Marti, M.D.
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/dbp/image/image.html
Insert Length: 1043 Std Error: 0.00
Seq primer: -40m13 fwd. EF from Amersham
High quality sequence stop: 336.
Location/Qualifiers
1. 395
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1369092"
/clone_11b="NCI-CGAP-GCB1"
/tissue_type="germinal center B cell"
/lab_host="DH10B"
/note="Vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was prepared from human tonsillar cells enriched for
germinal center B cells by flow sorting (CD20+, IgD-),
provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
(NCI) and Dr. Gerald Marti (CBER). CDNA synthesis was
primed with a Not I - oligo(dT) primer
[5'-TGTTACCAATCTGAAAGTGGAGCGCGCTCATTTTTTTTTTTTTTTT-3'

```

```

1. Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT7T3 vector. Library
went through one round of normalization, and was
constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT      100 a      77 c      73 g      145 t
ORIGIN

Query Match      83.6%; Score 18.4; DB 9; Length 395;
Best Local Similarity 95.0%; Pred. No. 4.3e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY      3 GACTTCTGAGTACCAATCAA 22
Db      345 GACTTCTGAGTACCAATCAA 326

RESULT 3
AA229942/c      364 bp      mRNA      linear      EST 21-AUG-1997
LOCUS      nc51h06.r1 NCI-CGAP_Pr3 Homo sapiens CDNA clone IMAGE:1011707
DEFINITION      similar to contains Alu repetitive element, mRNA sequence.
ACCESSION      AA229942
VERSION      AA229942.1 GI:1852255
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 364)
REFERENCE      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
TITLE      Tumor Gene Index
JOURNAL      Unpublished (1997)
COMMENT      Contact: Robert Strausberg, Ph.D.
Email: cgabs-remail.nih.gov
Tissue Procurement: W. Marston Linehan, M.D., Rodrigo Chuagui, M.D.
Michael Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: David B. Krizman, Ph.D.
CDNA Library Arrayed by: Genome Systems Inc., Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/dbp/image/image.html
Insert Length: 616 Std Error: 0.00
Seq primer: -28m13 rev1 EF from Amersham.
Location/Qualifiers
1. 364
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1011707"
/clone_11b="NCI-CGAP_Pr3"
/sex="Male"
/dev_stage="45 years old"
/lab_host="DH10B"
/note="Vector: PAMF10; Site_1: NotI; Site_2: EcoRI; 1st
strand cDNA was primed with oligo(dT)17 on 50 ng of
DNase-treated, total cellular RNA obtained from 5,000-10
,000 microdissected cells histologically-determined to be
fully malignant prostate cancer cells. Double-stranded
cDNA was ligated to EcoRI adaptors, 5 cycles of PCR
applied to the cDNA with an adaptor-specific primer, and
the resulting PCR product subcloned into PAMF10 by the
UDC-cloning method (Life Technologies). Average insert
size is 600 bp. NOTE: Not directionally cloned. This
library was constructed by David Krizman."

```

```

BASE COUNT      101 a      82 c      86 g      95 t
ORIGIN

Query Match      80.9%; Score 17.8; DB 9; Length 364;
Best Local Similarity 90.5%; Pred. No. 7.9e+02;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```



OY 2 AGACTTCTGAGTAACATCAA 22  
 Db 48 AGCTCTGCTGAGTAACATCAA 28

# RESULT 4

BO622226

LOCUS 418 bp mRNA linear EST 01-JUL-2002

DEFINITION fchlc.pk005.h2 Conidiobolus coronatus ARSEF 512 Conidiobolus

ACCESSION BO622226

VERSION BO622226.1 GI:21649395

KEYWORDS EST

SOURCE Conidiobolus coronatus

ORGANISM Conidiobolus coronatus

REFERENCE 1 (bases 1 to 418)

AUTHORS Fieldner, F.M., Screen, S., Hu, G., and St. Leger, R.J.

TITLE EST analysis of genes expressed by the zygomycete pathogen

JOURNAL Conidiobolus coronatus during optimized secretion of proteins

COMMENT Unpublished (2002)

Contact: Fieldner, F.M.

Department of Entomology

University of Maryland

4112 Plant Sciences Building, College Park, MD 20742, USA

Tel: 301 405 16 13

Fax: 301 314 92 90

Email: ffieldner@umd.edu

Location/Qualifiers

1. .418

/organism="Conidiobolus coronatus"

/strain="ARSEF 512"

/db\_xref="taxon:34488"

/clone\_id="Conidiobolus coronatus ARSEF 512"

/note="Vector: Unizap; Conidiobolus coronatus was grown in

minimal medium supplemented with Manduca sexta cuticle and

peptide for 18 hours. A cDNA library was constructed in

the unidirectional Lambda vector Unizap."

BASE COUNT 119 a 85 c 76 g 136 t

ORIGIN 2 others

Query Match 80.9%; Score 17.8; DB 14; Length 418;

Best Local Similarity 90.5%; Pred. No. 8.3e+02;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 AGACTTCTGAGTAACATCAA 22

Db 175 AGACTTCTGAGTAACATCAA 195

RESULT 5

LOCUS 509 bp mRNA linear EST 29-NOV-2001

DEFINITION sa94907.y1 Gm-cl065 glycine max cDNA clone GENOME SYSTEMS CLONE

ACCESSION BG882650

VERSION BG882650

KEYWORDS EST

SOURCE soybean

ORGANISM Glycine max

REFERENCE 1 (bases 1 to 509)

AUTHORS Shoemaker, R., Kelm, P., Vodkin, L., Erpelting, J., Coryell, V., Khanna

, A., Bolla, B., Merritt, M., Hillier, L., Kucaba, T., Martin, J., Beck, C.,

Wyllie, T., Underwood, K., Stepien, M., Theising, B., Allen, M., Bowers

, Y., Pearson, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk

, R., Ritter, E., Kohn, S., Sridh, T., Jackson, Y., Cardenas, M., McCann

, R., Waterston, R., and Wilson, R.

TITLE Public Soybean EST Project

# JOURNAL COMMENT

Unpublished (1999)  
 Contact: Shoemaker R/Public Soybean EST Project  
 Public Soybean EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: est@wustl.edu

# FEATURES

1. .509

# source

/organism="Glycine max"

# location/qualifiers

/db\_xref="taxon:3847"

# clone\_id

/clone\_id="Gm-cl065"

# issue\_type

/issue\_type="germinating shoots"

# lab\_host

/lab\_host="DH10B"

# note

/note="Vector: Bluescript II SK+; Site 1: EcoRI; Site 2: XhoI. The cDNA library was constructed from mRNA isolated

# germinating shoots

of the cultivar Williams. The seeds were allowed to germinate for 24 hours prior to being

# could stressed

for 2 days at 4C. Complementary DNA was synthesized from mRNA using a primer consisting of a

# poly(dT)

sequence with a XhoI restriction site. EcoRI adapters were ligated to the blunt-ended cDNA fragments

# directionally cloned

into the EcoRI-XhoI restriction site of the plasmid vector. The ligated cDNA fragments were

# transformed into

DH10B host cells (GibcoBRL). This library was constructed in the laboratory of Dr. Randy

# Shoemaker."

# BASE COUNT

151 a 91 c 121 g 146 t

# ORIGIN

# Query Match

80.9%; Score 17.8; DB 12; Length 509;

# Best Local Similarity

90.5%; Pred. No. 8.9e+02;

# Matches 19; Conservative

0; Mismatches 2; Indels 0; Gaps 0;

# OY 2

AGACTTCTGAGTAACATCAA 22

# Db 77

AGCTTCTGAGTAACATCAA 57

# RESULT 6

LOCUS 545 bp mRNA linear EST 20-DEC-1999

# DEFINITION

w122a08.x1 NCI-CGAP Col6 Homo sapiens cDNA clone IMAGE:2390966 3'

# ACCESSION

A1738689

# VERSION

A1738689

# KEYWORDS

EST

# SOURCE

human

# ORGANISM

Homo sapiens

# REFERENCE

1 (bases 1 to 545)

# AUTHORS

NCI-CGAP http://www.nci.nih.gov/ncicgap

# JOURNAL

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

# COMMENT

Unpublished (1997)

# Contact:

Robert Strussberg, Ph.D.

# Email:

cgapb@remail.nih.gov

# Tissue Procurement:

Ilan Kirsch, M.D., Michael R. Emmert-Buck, M.D.

# , Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

# cDNA Library Arrayed by:

Greg Lennon, Ph.D.

# DNA Sequencing by:

Washington University Genome Sequencing Center

# Clone distribution:

NCI-CGAP clone distribution information can be

# found through the

I.M.A.G.E. Consortium/LLNL at:

# www.bio.lnl.gov

db/rp/image/image.html



Db 509 AAGACTCTTGTAACAATCA 489

RESULT 9  
LOCUS B609955  
DEFINITION NXS1\_053\_A10\_F NXS1 (Nsf Xylem Side wood Inclined) Pinus taeda cDNA  
clone NXS1\_053\_A10 5', mRNA sequence.  
ACCESSION B609955  
VERSION B609955.1 GI:11778131  
KEYWORDS EST.  
SOURCE loblolly pine.  
ORGANISM Pinus taeda  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Coniferopsida; Coniferales; Pinaceae; Pinus; Pinus.  
REFERENCE 1 (bases 1 to 375)  
AUTHORS Sederoff, R.  
TITLE Molecular Basis of Wood Formation in the Pine Megagenome  
JOURNAL Unpublished (2000)  
COMMENT Contact: Johnson, Arthur  
North Carolina State University  
Tel: 919 515 7800  
Fax: 919 515 7801  
Email: rjohnson@unity.ncsu.edu  
Seq primer: 73.  
Location/Qualifiers  
1..375  
/organism="Pinus taeda"  
/strain="Coastal plain loblolly pine from North Carolina"  
/db\_xref="taxon:3352"  
/clone="NXS1\_053\_A10"  
/clone\_1lb="NXS1 (Nsf Xylem Side wood Inclined)"  
/tissue\_type="xylem"  
/cell\_type="Side"  
/dev\_stage="Juvenile"  
/lab\_host="XLI-Blue"  
/note="Vector: Bluescript SK; Site\_1: Eco RI; Site\_2: Xho I  
: The library is from early (spring) wood, taken from  
three six-year old trees (three different genotypes), in  
the juvenile phase. These trees were induced to form side  
wood by bending to a 45 degree angle and tying them to the  
ground. Differentiating xylem was harvested from the sides  
of the inclined stems, and a mixture of all three  
genotypes was used for the library. oligo-dt primed cDNA  
was directionally cloned into the EcoRI-XhoI Bluescript SK  
vector arms. NOTE: The sequences contain a 'cDNA adapter'  
between the EcoRI site and the start of the EST. The  
adapter sequence is 'AATTCGACGAG'."

BASE COUNT 80 a 76 c 76 g 129 t 14 others  
ORIGIN

Query Match 79.1%; Score 17.4; DB 12; Length 375;  
Best Local Similarity 94.7%; Pred. No. 1.2e+03;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 ACTCTGAGTACATCA 22  
|||||  
Db 132 ACTCTGAGTACATCA 150

RESULT 10  
LOCUS AM699516/c  
DEFINITION 474 bp mRNA linear EST 18-APR-2000  
9db6f02.y1 Moss EST library PPN Physcomitrella patens cDNA clone  
PEP\_SOURCE\_ID:PPN080304 5' similar to TR:096538 Q96538 ACYL-COA  
SYNTHETASE; mRNA sequence.  
ACCESSION AM699516  
VERSION AM699516.1 GI:7583607  
KEYWORDS EST.  
SOURCE Physcomitrella patens.  
ORGANISM Physcomitrella patens  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Bryophyta;  
Bryopsida; Funariidae; Funariales; Funariaceae; Physcomitrella.

REFERENCE 1 (bases 1 to 474)  
AUTHORS Quatrano, R., Bashardes, S., Cove, D., Cumming, A., Knight, C., Clifton  
Marra, M., Hillier, L., Page, D., Martin, J., Wylie, T., Underwood  
R., Theising, B., Allen, M., Bowers, Y., Person, B., Swaller, T.,  
Steele, M., Gibbons, M., Harvey, N., Rafter, E., Jackson, Y., McCann, R.,  
Waterson, R., and Wilson, R.  
TITLE Leeds/Mash U Moss EST Project  
JOURNAL Unpublished (1999)  
COMMENT Contact: Ralph Quatrano  
Leeds/Mash U Moss EST Project  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: est@watson.wustl.edu  
Libraries were constructed by Dr. Stavros Bashardes as part of the  
Physcomitrella EST program (PEP) at the Univ. of Leeds (UK) and  
Washington Univ. in St. Louis (USA) DNA sequencing by: Washington  
University Genome Sequencing Center For information on obtaining a  
clone please contact: Celia Knight (c.d.knight@leeds.ac.uk)  
Seq primer: -40RP from Gibco  
High quality sequence stop: 405.  
Location/Qualifiers  
1..474  
/organism="Physcomitrella patens"  
/db\_xref="taxon:3218"  
/clone="PEP\_SOURCE\_ID:PPN080304"  
/clone\_1lb="Moss EST library PPN"  
/tissue\_type="protonemata: 7 day old tissue auxin treated"  
/lab\_host="DH10B"  
/note="Vector: Bluescript SK; Site\_1: EcoRI; Site\_2:  
XhoI: Construction of the cDNA library was carried out  
using Stratagene's 'Unizap - cDNA synthesis kit'. cDNA was  
constructed using an oligo dt primer/linker that contains  
a XhoI site within it. Following ds cDNA synthesis,  
EcoRI adapters were ligated to the blunt ends and sample  
sticky end on one side and a XhoI sticky end on the other.  
This cDNA was ligated directionally in Unizap arms. The  
vector is designed containing the pluescript sequence as  
well as lambda DNA and cDNA is cloned within this  
pluescript sequence. The vector was then packaged using  
Gold gigacloning extracts. Library was grown in XL1blue  
MRP cells and amplified. The library was excised by mass  
excision using Stratagene's 'Mass excision kit' that uses  
exsist as a helper phage that releases the pluescript  
sequence and circularises it as single stranded plasmids  
that are then packaged (by helper phage) and secreted out  
of the host cell as phagemids. SOLR cells were transformed  
with phagemids and the library was plated out on LB-amp  
plates to select for transformants. Approximately 1,000  
,000 colonies were grown and recovered. The double  
stranded plasmid library was recovered by using Qulagen  
Midl prep kit. 2 micro grams of each library were used to  
transform DH10B cells by electroporation."

BASE COUNT 138 a 99 c 114 g 123 t  
ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 474;  
Best Local Similarity 94.7%; Pred. No. 1.3e+03;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 ACTCTGAGTACATCA 22  
|||||  
Db 432 ACTCTGAGTACATCA 414

RESULT 11  
LOCUS BG040453  
DEFINITION 477 bp mRNA linear EST 24-JAN-2001  
NXS1\_108\_D07\_F NXS1 (Nsf Xylem Side wood Inclined) Pinus taeda cDNA  
clone NXS1\_108\_D07 5', mRNA sequence.  
ACCESSION BG040453

VERSION BG040453.1 GI:12483038  
 KEYWORDS EST.  
 SOURCE loblolly pine.  
 ORGANISM Pinus taeda

REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermotophyta; Coniferopsida; Coniferales; Pinaceae; Pinus.  
 TITLE 1 (bases 1 to 477)  
 AUTHORS Sederoff, R.  
 JOURNAL Molecular Basis of Wood Formation in the Pine Megagenome  
 COMMENT Unpublished (2000)  
 Contact: Johnson, Arthur  
 North Carolina State University  
 Tel: 919 515 7800  
 Fax: 919 515 7801  
 Email: ajohnson@unity.ncsu.edu  
 Seq primer: T3

FEATURES  
 source  
 1..477  
 /organism="Pinus taeda"  
 /strain="Coastal plain loblolly pine from North Carolina"  
 /db\_xref="taxon:3352"  
 /clone="NXSL108.D07"  
 /clone\_lib="NXSL (Nsf Xylem Side wood Inclined)"  
 /tissue\_type="Xylem"  
 /cell\_type="Side"  
 /dev\_stage="juvenile"  
 /lab\_host="XLI-Blue"  
 /note="Vector: Bluescript SK; Site.1: Eco RI; Site.2: Xho I  
 ; The library is from early (spring) wood, taken from  
 three six-year old trees (three different genotypes), in  
 the juvenile phase. These trees were induced to form side  
 wood by bending to a 45 degree angle and tying them to the  
 ground. Differentiating xylem was harvested from the sides  
 of the inclined stems, and a mixture of all three  
 genotypes was used for the library. oligo-dT primed cDNA  
 was directionally cloned into the EcoRI-XhoI Bluescript SK  
 vector arms. NOTE: The sequences contain a 'cDNA adapter'  
 between the EcoRI site and the start of the EST. The  
 adapter sequence is 'ATTGCGACGACG'."

BASE COUNT 106 a 106 c 93 g 154 t 18 others

ORIGIN

Query Match 79.1%; Score 17.4; DB 12; Length 477;  
 Best Local Similarity 94.7%; Pred. No. 1.3e+03;  
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 ACTTCGAGTAACATCAA 22  
 ||||||||||||||||  
 Db 134 ACTTCGAGTAACATCAA 152

RESULT 12  
 AM738913 549 bp mRNA linear EST 25-APR-2000  
 AM738913/c GB1606.y1 Moss EST library PPN Physcomitrella patens cDNA clone  
 LOCUS PEP\_SOURCE\_ID:PPN100912 5' similar to TR:Q96538 Q96538 ACTL-COA  
 DEFINITION SYNTHETASE ; mRNA sequence.  
 ACCESSION AM738913  
 VERSION AM738913.1 GI:7647930  
 KEYWORDS EST.  
 SOURCE Physcomitrella patens.  
 ORGANISM Physcomitrella patens  
 REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Bryophyta;  
 Bryopsida; Funariidae; Funariales; Funariaceae; Physcomitrella.  
 TITLE 1 (bases 1 to 549)  
 AUTHORS Quatrano, R., Bashlars, S., Cove, D., Cuming, A., Knight, C., Clifton  
 S., Maitra, M., Hillier, L., Pape, D., Martin, U., Wylie, T., Underwood  
 K., Theising, B., Allen, M., Bowers, Y., Person, B., Swaller, T.,  
 Stepoe, M., Gibbons, M., Harvey, N., Ritzer, E., Jackson, Y., McCann, R.,  
 Waterston, R. and Wilson, R.  
 Leeds/Mash U Moss EST Project  
 COMMENT Unpublished (1999)  
 Contact: Ralph Quatrano

Leeds/Mash U Moss EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: est@wustl.edu

Libraries were constructed by Dr. Stavros Bashlars as part of the  
 Physcomitrella EST program (PEP) at the Univ. of Leeds (UK) and  
 Washington Univ. in St. Louis (USA) DNA sequencing by: Washington  
 University Genome Sequencing Center For information on obtaining a  
 clone please contact: Celia Knight (c.d.knight@leeds.ac.uk)  
 Seq primer: -40RP from Glibco  
 High quality sequence stop: 408.

FEATURES  
 source  
 1..549  
 /organism="Physcomitrella patens"  
 /db\_xref="taxon:3218"  
 /clone="PEP\_SOURCE\_ID:PPN100912"  
 /clone\_lib="Moss EST library PPN"  
 /tissue\_type="Protonemata: 7 day old tissue auxin treated"  
 /lab\_host="DH10B"  
 /note="Vector: pBluescript SK-; Site.1: EcoRI; Site.2:  
 XhoI; Construction of the cDNA library was carried out  
 using Stratagene's 'Unizap - cDNA synthesis kit'. cDNA was  
 constructed using an oligo-dT primer/linker that contains  
 a XhoI site within it. Following ds cDNA synthesis,  
 EcoRI adapters were ligated to the blunt ends and sample  
 was digested with XhoI. The result is cDNA with an EcoRI  
 sticky end on one side and a XhoI sticky end on the other.  
 This cDNA was ligated directionally in Unizap arms. The  
 vector is designed containing the pBluescript sequence as  
 well as lambda DNA and cDNA is cloned within this  
 pBluescript sequence. The vector was then packaged using  
 Gold gigapacking extracts. Library was grown in XL1Blue  
 MRF' cells and amplified. The library was excised by mass  
 excision using Stratagene's 'Mass excision kit' that uses  
 exasist as a helper phage that releases the pluscript  
 sequence and circularises it as single stranded plasmids  
 that are then packaged (by helper phage) and secreted out  
 of the host cell as phagemids. SOLR cells were transformed  
 with phagemids and the library was plated out on LB-amp  
 plates to select for transformants. Approximately 1,000  
 ,000 colonies were grown and recovered. The double  
 stranded plasmid library was recovered by using Qulagen  
 Midi prep kit. 2 micro grams of each library were used to  
 transform DH10B cells by electroporation."

BASE COUNT 151 a 116 c 125 g 157 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 549;  
 Best Local Similarity 94.7%; Pred. No. 1.4e+03;  
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 ACTTCGAGTAACATCAA 22  
 ||||||||||||||||  
 Db 418 ACTTCGAGTAACATCAA 400

RESULT 13  
 AO525332 608 bp DNA linear GSS 11-MAY-1999  
 LOCUS HS\_5226\_B1\_C08\_T7A RPT-11 Human Male BAC Library Homo sapiens  
 DEFINITION genomic clone Plate=802 COL=15 Row=F, DNA sequence.  
 ACCESSION AO525332  
 VERSION AO525332.1 GI:4772652  
 KEYWORDS GSS.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 TITLE 1 (bases 1 to 608)  
 AUTHORS Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T.,  
 Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and

**TITLE** Hood, L.  
Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

**JOURNAL** Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

**MEDLINE** 99380589

**COMMENT** Contact: Mahatras GG, Wallace JC, Hood L  
High Throughput Sequencing Center  
University of Washington  
401 Queen Anne Avenue North, Seattle, WA 98109, USA  
Tel: (206) 616-3618  
Fax: (206) 616-3687  
Email: jwallace@u.washington.edu

**FEATURES** Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@jlong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering\_bac.htm) or from Research Genetics (info@resgen.com). BAC end Web Server: http://www.hsc.washington.edu  
Plate: 802 row: F column: 15  
Seq primer: 77  
Class: BAC ends  
High quality sequence stop: 608.

**SOURCE** Location/Qualifiers

1..608  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="Plate-802 COL-15 Row-F"  
/clone\_lib="RPCI-11 Human Male BAC Library"  
/sex="male"

**NOTE** Vector: pBAC3.6; Site\_1: EcoRI; Site\_2: EcoRI;  
Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBAC3.6 vector at EcoRI sites 10 others

**BASE COUNT** 221 a 123 c 107 g 147 t 10 others

**Query Match** 79.1%; Score 17.4; DB 17; Length 608;  
Best Local Similarity 85.7%; Pred. No. 1.4e+03;  
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

**OY** 1 AAGCTTGAGTACATCA 21  
|||||  
Db 22 AAGCTTTTACATCA 42

**RESULT 14** Buj173371 611 bp mRNA linear EST 24-JAN-2002  
Buj173371 full length cDNA library, chloronemata and young gametophores Physcomitrella patens subsp. patens cDNA clone pph35912 3', mRNA sequence.

**ACCESSION** Buj173371 GI:18341336

**VERSION** EST.

**KEYWORDS** Physcomitrella patens subsp. patens.

**SOURCE** Physcomitrella patens subsp. patens  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Bryophyta; Bryopsida; Funariidae; Funariales; Funariaceae; Physcomitrella.

**ORGANISM** 1 (bases 1 to 611)  
Fujita, T., Shin-I, T., Seki, M., Kamiya, A., Uchiyama, I., Nishiyama, T., Carninci, P., Hayashizaki, Y., Shinozaki, K., Kohara, Y. and Hasebe

**REFERENCE** 'M. Carninci, P., Hayashizaki, Y., Shinozaki, K., Kohara, Y. and Hasebe

**AUTHORS** 'M. Carninci, P., Hayashizaki, Y., Shinozaki, K., Kohara, Y. and Hasebe

**TITLE** Comparison of the moss Physcomitrella patens genome with flowering plants genome

**JOURNAL** Unpublished (2002)

**COMMENT** Contact: Tadasu Shin-I  
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A backbone of the vector is basically from pBluescript(KS), that was in vivo excised from a modified IPS phage vector (Mo bi rec, Germany). 5' end of the cDNA that was digested with XhoI was ligated to SalI site of the vector and the 3' end including polyA tail was ligated to BamHI site of the vector. cDNA insert could be amplified with conventional T7 and T3 primers. This full-length cDNA library was generated basically according to the method described in The Plant J 15, 707-720 (1998) Seki M. et al. Protonemata were blended by the POLYTRON, and then cultivated on the BCDATG medium for 13-14 days under the continuous light.

**FEATURES** Location/Qualifiers

1..611

/organism="Physcomitrella patens subsp. patens"  
/db\_xref="taxon:145481"  
/clone="pph35912"  
/clone\_lib="full length cDNA library, chloronemata and young gametophores"  
/tissue\_type="mixture of chloronemata and young gametophores with 2 to 5 leaves"

**BASE COUNT** 178 a 135 c 128 g 170 t

**ORIGIN**

**Query Match** 79.1%; Score 17.4; DB 13; Length 611;  
Best Local Similarity 94.7%; Pred. No. 1.4e+03;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

**OY** 4 ACTTCTGAGTACATCA 22  
|||||  
Db 181 ACTTCTGAGTACATCA 199

**RESULT 15** Buj171523 692 bp mRNA linear EST 24-JAN-2002  
Buj171523 full length cDNA library, chloronemata and young gametophores Physcomitrella patens subsp. patens cDNA clone pph29d18 3', mRNA sequence.

**ACCESSION** Buj171523 GI:18339496

**VERSION** EST.

**KEYWORDS** Physcomitrella patens subsp. patens.

**SOURCE** Physcomitrella patens subsp. patens  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Bryophyta; Bryopsida; Funariidae; Funariales; Funariaceae; Physcomitrella.

**ORGANISM** 1 (bases 1 to 692)  
Fujita, T., Shin-I, T., Seki, M., Kamiya, A., Uchiyama, I., Nishiyama, T., Carninci, P., Hayashizaki, Y., Shinozaki, K., Kohara, Y. and Hasebe

**REFERENCE** 'M. Carninci, P., Hayashizaki, Y., Shinozaki, K., Kohara, Y. and Hasebe

**AUTHORS** 'M. Carninci, P., Hayashizaki, Y., Shinozaki, K., Kohara, Y. and Hasebe

**TITLE** Comparison of the moss Physcomitrella patens genome with flowering plants genome

**JOURNAL** Unpublished (2002)

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A backbone of the vector is basically from pBluescript(KS), that was in vivo excised from a modified IPS phage vector (Mo bi rec, Germany). 5' end of the cDNA that was digested with XhoI was ligated to SalI site of the vector and the 3' end including polyA tail was ligated to BamHI site of the vector. cDNA insert could be amplified with conventional T7 and T3 primers. This full-length cDNA library was generated basically according to the method described in The Plant J 15, 707-720 (1998) Seki M. et al. Protonemata were blended by the POLYTRON, and then cultivated on the BCDATG medium for 13-14 days under the continuous light.

**FEATURES** Location/Qualifiers

1..692

/organism="Physcomitrella patens subsp. patens"  
/db\_xref="taxon:145481"  
/clone="pph29d18"

/clone.lib="full length cDNA library, chloronemata and  
young gametophores"  
/tissue.type="mixture of chloronemata and young  
gametophores with 2 to 5 leaves"  
BASE COUNT 213 a 153 c 148 g 178 t  
ORIGIN

Query Match 79.1%; Score 17.4; DB 13; Length 692;  
Best Local Similarity 94.7%; Pred. No. 1.5e+03;  
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 AAGACTTCTGAGTACAAAT 19  
|||||  
Db 337 AAGACTTCTGAGTACAAAT 355

Search completed: March 17, 2003, 13:09:28  
Job time : 782.688 secs